PCT WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 :
C07D 401/12, 401/06, 211/14, 211/22, A61K 31/445, A61P 11/00, 19/00, 31/00

A1

(11) International Publication Number: WO 00/58305

(43) International Publication Date: 5 October 2000 (05.10.00)

(21) International Application Number: PCT/SE00/00563

(22) International Filing Date: 22 March 2000 (22.03.00)

(30) Priority Data:

9901117-3	26 March 1999 (26.03.99)	SE
9902194-1	10 June 1999 (10.06.99)	SE

(71) Applicant *(for all designated States except MG US)*: ASTRAZENECA UK LIMITED [GB/GB]; 15 Stanhope Gate, London W1Y 6LN (GB).

(71) Applicant *(for MG only)*: ASTRAZENECA AB [SE/SE]; S-151 85 Södertälje (SE).

(72) Inventors; and

(75) Inventors/Applicants *(for US only)*: BAXTER, Andrew [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB). BROUGH, Stephen [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB). KINDON, Nicholas [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB). MCINALLY, Thomas [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB). ROBERTS, Bryan [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB). THOM, Stephen [GB/GB]; AstraZeneca, R & D Charnwood, Bakewell Road, Loughborough, Leisc. LE11 5RH (GB).

(74) Agent: ASTRAZENECA AB; Global Intellectual Property, Patents, S-151 85 Södertälje (SE).

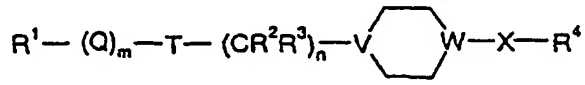
(81) Designated States: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published

With international search report.

Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: NOVEL COMPOUNDS



(I)

(57) Abstract

The invention provides compounds of general formula (I) wherein: R^1 represents optionally substituted C_1–C_{12} alkyl or optionally substituted 3- to 10-membered saturated or unsaturated ring system comprising up to two ring carbon atoms that form carbonyl groups and comprising up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur; m is 0–1; Q represents OCH_2, C_1–C_4 alkylene or C_2–C_4 alkenylene; T represents C(O)NH, or when m is 0, T may additionally represent a bond or NH, or when m is 1 and Q represents C_1–C_4 alkylene, T may additionally represent NH; n is 1–4; each R^2 and R^3 independently represents H or C_1–C_4 alkyl; V represents N, and W represents N or CH; X represents O, C(O), CH(OH), SO_2, NH or N(C_1–C_6 alkyl), provided that when W represents N, then X represents either C(O) or SO_2 and when W represents CH, then X is other than SO_2; R^4 represents optionally substituted phenyl; R^5 and R^6 each independently represent H, C_1–C_6 alkyl or hydroxyC_1–C_6 alkyl, or R^5 and R^6 together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring; R^7 and R^8 each independently represent H or C_1–C_6 alkyl; and R^9 represents OH or $-NR^7R^8$; processes for their preparation, pharmaceutical compositions containing them and their use in therapy.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the P

AL	Albania
AM	Armenia
AT	Austria
AU	Australia
AZ	Azerbaijan
BA	Bosnia and Herzegovina
BB	Barbados
BE	Belgium
BF	Burkina Faso
BG	Bulgaria
BJ	Benin
BR	Brazil
BY	Belarus
CA	Canada
CF	Central African Republic
CG	Congo
CH	Switzerland
CI	Côte d'Ivoire
CM	Cameroon
CN	China
CU	Cuba
CZ	Czech Republic
DE	Germany
DK	Denmark
EE	Estonia
ES	Spain
FI	Finland
FR	France
GA	Gabon
GB	United Kingdom
GE	Georgia
GH	Ghana
GN	Guinea
GR	Greece
HU	Hungary
IE	Ireland
IL	Israel
IS	Iceland
IT	Italy
JP	Japan
KE	Kenya
KG	Kyrgyzstan
KP	Democratic People's Republic of Korea
KR	Republic of Korea
KZ	Kazakstan
LC	Saint Lucia
LI	Liechtenstein
LK	Sri Lanka
LR	Liberia
LS	Lesotho
LT	Lithuania
LU	Luxembourg
LV	Larvia
MC	Monaco
MD	Republic of Moldova
MG	Madagascar
MK	The former Yugoslav Republic of Macedonia
ML	Mali
MN	Mongolia
MR	Mauritania
MW	Malawi
MX	Mexico
NE	Niger
NL	Netherlands
NO	Norway
NZ	New Zealand
PL	Poland
PT	Portugal
RO	Romania
RU	Russian Federation
SD	Sudan
SE	Sweden
SG	Singapore
SI	Slovenia
SK	Slovakia
SN	Senegal
SZ	Swaziland
TD	Chad
TG	Togo
TJ	Tajikistan
TM	Turkmenistan
TR	Turkey
TT	Trinidad and Tobago
UA	Ukraine
UG	Uganda
US	United States of America
UZ	Uzbekistan
VN	Viet Nam
YU	Yugoslavia
ZW	Zimbabwe

NOVEL COMPOUNDS

The present invention relates to novel compounds, processes for their preparation, pharmaceutical compositions containing them and their use in therapy.

Chemokines play an important role in immune and inflammatory responses in various diseases and disorders, including asthma and allergic diseases, as well as autoimmune pathologies such as rheumatoid arthritis and atherosclerosis. These small secreted molecules are a growing superfamily of 8-14 kDa proteins characterised by a conserved four cysteine motif. The chemokine superfamily can be divided into two main groups exhibiting characteristic structural motifs, the Cys-X-Cys (C-X-C) and Cys-Cys (C-C) families. These are distinguished on the basis of a single amino acid insertion between the NH-proximal pair of cysteine residues and sequence similarity.

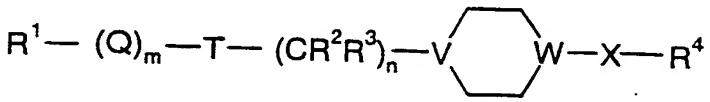
The C-X-C chemokines include several potent chemoattractants and activators of neutrophils such as interleukin-8 (IL-8) and neutrophil-activating peptide 2 (NAP-2).

The C-C chemokines include potent chemoattractants of monocytes and lymphocytes but not neutrophils such as human monocyte chemotactic proteins 1-3 (MCP-1, MCP-2 and MCP-3), RANTES (Regulated on Activation, Normal T Expressed and Secreted), eotaxin and the macrophage inflammatory proteins 1α and 1β (MIP-1α and MIP-1β).

Studies have demonstrated that the actions of the chemokines are mediated by subfamilies of G protein-coupled receptors, among which are the receptors designated CCR1, CCR2, CCR2A, CCR2B, CCR3, CCR4, CCR5, CCR6, CCR7, CCR8, CCR9, CCR10, CXCR1, CXCR2, CXCR3 and CXCR4. These receptors represent good targets for drug development since agents which modulate these receptors would be useful in the treatment of disorders and diseases such as those mentioned above.

Certain piperidinyl derivatives and piperazinyl derivatives are known from U.S. Patents Nos. 3 787 419, 4 559 349 and 5 210 086 for use respectively as central nervous system depressants, antipsychotic agents and as α_1-adrenoreceptor antagonists.

In accordance with the present invention, there is therefore provided a compound of general formula



(I)

wherein:

R^1 represents a C_1-C_{12} alkyl group optionally substituted by one or more substituents independently selected from cyano, hydroxyl, C_1-C_6 alkoxy, C_1-C_6 alkylthio and C_1-C_6 alkoxycarbonyl groups, or

R^1 represents a 3- to 10-membered saturated or unsaturated ring system which may comprise up to two ring carbon atoms that form carbonyl groups and which may comprise up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur, the ring system being optionally substituted by one or more substituents independently selected from halogen atoms, and cyano, nitro, hydroxyl, C_1-C_6 alkyl, C_3-C_6 cycloalkyl, C_1-C_6 alkoxy, C_1-C_6 alkoxycarbonyl, C_1-C_6 haloalkyl, C_1-C_6 haloalkoxy, $-NR^5R^6$, C_3-C_6 cycloalkylamino, C_1-C_6 alkylthio, C_1-C_6 alkylthioC_1-C_6 alkyl, C_1-C_6 alkylcarbonylamino, $-C(O)NR^7R^8$, sulphonamido ($-SO_2NH_2$), (di)C_1-C_6 alkylsulphonamido, phenyl, phenylamino, nitrophenyl, pyridyl, pyridylthio, benzodioxanyl, thienyl, furanyl, and $C(O)R^9$-substituted C_1-C_6 alkyl or C_1-C_6 alkoxy groups;

m is 0 or 1;

Q represents a group OCH_2, C_1-C_4 alkylene or C_2-C_4 alkenylene;

T represents a group C(O)NH, or when m is 0, T may additionally represent a bond or a group NH, or when m is 1 and Q represents C_1-C_4 alkylene, T may additionally represent a group NH;

n is 1, 2, 3 or 4;

each R^2 independently represents a hydrogen atom or a C_1-C_4 alkyl group;

each R^3 independently represents a hydrogen atom or a C_1-C_4 alkyl group;

V represents a nitrogen atom;

W represents a nitrogen atom or a group CH;

X represents an oxygen atom or a group C(O), CH(OH), NH or N(C_1-C_6 alkyl), provided that when W represents a nitrogen atom, then X represents C(O);

R^4 represents a phenyl group optionally substituted by one or more substituents independently selected from halogen atoms, and amino, nitro, cyano, sulphonyl (-SO_3H), sulphonamido (-SO_2NH_2), C_1-C_6 alkyl, C_1-C_6 haloalkyl, C_1-C_6 haloalkoxy and C_1-C_6 alkylsulphonyl groups;

R^5 and R^6 each independently represent a hydrogen atom or a C_1-C_6 alkyl or hydroxyC_1-C_6 alkyl group, or R^5 and R^6 together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring;

R^7 and R^8 each independently represent a hydrogen atom or a C_1-C_6 alkyl group; and

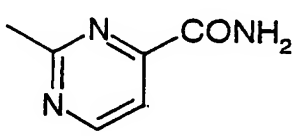
R^9 represents a hydroxyl or -NR^7R^8 group;

with the provisos that

(a) when m is 0, T is CONH, n is 2, 3 or 4 and each R^2 and R^3 represents hydrogen, W is CH, X is C(O) or CH(OH) and R^1 represents a substituted 3- to 10-membered unsaturated ring system, then the one or more substituents in the ring system do not include hydroxyl, halogen, C_1-C_6 alkoxy or C_1-C_6 haloalkoxy, and

(b) when W is N, X is C(O), R^4 represents 3-trifluoromethylphenyl, m is 0 and T is a bond, then R^1 and $(CR^2R^3)_n$ taken together do not represent a C_1-C_6 alkyl group, and

(c) when W is CH, X is O, n is 2 or 3 and each R^2 and R^3 represents hydrogen, m is 0 and T is NH, then R^1 does not represent a group



;

or a pharmaceutically acceptable salt or solvate thereof.

In the context of the present specification, an alkyl substituent group or an alkyl moiety in a substituent group may be linear or branched. Further, the alkyl moieties in a dialkylamino, di(hydroxyalkyl)amino or dialkylsulphonamido substituent group may be the same or different.

R^1 represents a C_1-C_{12}, preferably C_1-C_{10}, alkyl group optionally substituted by one or more (e.g. one, two, three or four) substituents independently selected from cyano, hydroxyl, C_1-C_6, preferably C_1-C_4, alkoxy (e.g. methoxy, ethoxy, propoxy, butoxy, pentoxy or hexoxy), C_1-C_6, preferably C_1-C_4, alkylthio (e.g. methyl-, ethyl-, propyl-, butyl-, pentyl- or hexylthio) and C_1-C_6, preferably C_1-C_4, alkoxycarbonyl (e.g. methoxy-, ethoxy-, propoxy-, butoxy-, pentoxy- or hexoxycarbonyl) groups, or
R^1 represents a 3- to 10-membered saturated or unsaturated ring system comprising up to two ring carbon atoms that form carbonyl groups and comprising up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur, the ring system being optionally substituted by one or more (e.g. one, two, three or four) substituents independently selected from halogen atoms (fluorine, chlorine, bromine or iodine), and cyano, nitro, hydroxyl, C_1-C_6 alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl or hexyl), C_3-C_6 cycloalkyl (cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl), C_1-C_6 alkoxy (e.g. methoxy, ethoxy, propoxy, butoxy, pentoxy or hexoxy), C_1-C_6 alkoxycarbonyl (e.g. methoxy-, ethoxy-, propoxy-, butoxy-, pentoxy- or hexoxycarbonyl), C_1-C_6 haloalkyl (e.g. trifluoromethyl), C_1-C_6 haloalkoxy (e.g. trifluoromethoxy), $-NR^5R^6$, C_3-C_6 cycloalkylamino (cyclopropyl-, cyclobutyl-, cyclopentyl- or cyclohexylamino), C_1-C_6 alkylthio (e.g. methyl-, ethyl-, propyl-, butyl-, pentyl- or hexylthio), C_1-C_6 alkylthioC_1-C_6 alkyl (e.g. methylthiomethyl), C_1-C_6 alkylcarbonylamino (e.g. methyl-, ethyl-, propyl-, butyl-, pentyl- or hexylcarbonylamino), $-C(O)NR^7R^8$, sulphonamido ($-SO_2NH_2$), (di)C_1-C_6 alkylsulphonamido (e.g. (di)methylsulphonamido or (di)ethylsulphonamido), phenyl, phenylamino, nitrophenyl, pyridyl, pyridylthio, benzodioxanyl, thienyl,

furanyl, and $C(O)R^9$-substituted C_1-C_6 alkyl or C_1-C_6 alkoxy groups, the alkyl and alkoxy moieties being as defined above.

The 3- to 10-membered saturated or unsaturated ring system in the group R^1 may be monocyclic, or polycyclic comprising 2 or more fused rings, examples of which include cyclobutyl, cyclopentyl, cyclohexyl, norbornylenyl, adamantyl, piperidyl, phenyl, naphthyl, naphthyridinyl, 1,3-benzodioxolyl, pyrazolyl, furanyl, pyridyl, thienyl, benzoxazolyl, benzothiazolyl, chromonyl, imidazolyl, quinolinyl, isoquinolinyl, benzimidazolyl, pyrimidinyl, pyrazolopyrimidinyl, thienopyrimidinyl, thiazolopyrimidinyl, pyrimidinedione, pyrazinyl, pyridazinyl, purinyl, quinoxalinyl, thiazolyl, isothiazolyl and 2,4-dioxo-3,4-dihydro-quinazolinyl.

Preferably, R^1 represents a C_1-C_{10} alkyl group optionally substituted by one or two substituents independently selected from cyano, hydroxyl, C_1-C_4 alkoxy, C_1-C_4 alkylthio and C_1-C_4 alkoxycarbonyl groups, or

R^1 represents a 3- to 10-membered saturated or unsaturated ring system comprising up to two ring carbon atoms that form carbonyl groups and comprising up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur, the ring system being optionally substituted by one, two or three substituents independently selected from halogen atoms, and cyano, nitro, hydroxyl, C_1-C_4 alkyl, C_3-C_6 cycloalkyl, C_1-C_4 alkoxy, C_1-C_4 alkoxycarbonyl, C_1-C_3 haloalkyl, C_1-C_3 haloalkoxy, $-NR^5R^6$, C_3-C_6 cycloalkylamino, C_1-C_4 alkylthio, C_1-C_4 alkylthioC_1-C_4 alkyl, C_1-C_4 alkylcarbonylamino, $-C(O)NR^7R^8$, phenyl, phenylamino, nitrophenyl, pyridyl, pyridylthio, benzodioxanyl, thienyl, furanyl, and $C(O)R^9$-substituted C_1-C_4 alkyl or C_1-C_4 alkoxy groups.

Preferably Q represents a group OCH_2, C_1-C_3 alkylene or C_2-C_3 alkenylene.

Each R^2 independently represents a hydrogen atom or a C_1-C_4 alkyl (e.g. methyl, ethyl, propyl, isopropyl or butyl) group, and is especially a hydrogen atom.

Each R^3 independently represents a hydrogen atom or a C_1-C_4 alkyl (e.g. methyl, ethyl, propyl, isopropyl or butyl) group, and is especially a hydrogen atom.

Preferably n is 2 or 3.

X preferably represents an oxygen atom or a group C(O) or NH.

R^4 represents a phenyl group optionally substituted by one or more (e.g. one, two, three or four) substituents independently selected from halogen atoms (fluorine, cholorine, bromine or iodine), and amino, nitro, cyano, sulphonyl ($-SO_3H$), sulphonamido ($-SO_2NH_2$), C_1-C_6, preferably C_1-C_4, alkyl (e.g. methyl, ethyl, propyl, butyl, pentyl or hexyl), C_1-C_6, preferably C_1-C_4, haloalkyl (e.g. trifluoromethyl), C_1-C_6, preferably C_1-C_4, haloalkoxy (e.g. trifluoromethoxy) and C_1-C_6, preferably C_1-C_4, alkylsulphonyl (e.g. methyl-, ethyl-, propyl-, butyl-, pentyl- or hexylsulphonyl) groups.

Preferably, R^4 represents a phenyl group optionally substituted by one or two halogen atoms, particularly chlorine atoms.

R^5 and R^6 each independently represent a hydrogen atom or a C_1-C_6, preferably C_1-C_4, alkyl or hydroxyC_1-C_6, preferably C_1-C_4, alkyl group, or R^5 and R^6 together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring. The alkyl moiety in each case may, for example, be a methyl, ethyl, propyl, butyl, pentyl or hexyl group. In the hydroxyalkyl group, the hydroxyl group may be attached to any suitable carbon atom of the alkyl moiety.

R^7 and R^8 each independently represent a hydrogen atom or a C_1-C_6, preferably C_1-C_4, alkyl (e.g. methyl, ethyl, propyl, butyl, pentyl or hexyl) group. Preferably, R^7 and R^8 each independently represent a hydrogen atom or a methyl group.

R^9 represents a hydroxyl or, preferably, $-NR^5R^6$ group.

Examples of particularly preferred compounds of the invention include:
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-ethoxybenzamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-isopropoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-ethoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethoxy)benzamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride,
3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide hydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-hydroxybenzamide hydrochloride,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-[2-(methylamino)-2-oxoethoxy]benzamide hydrochloride,
2-[3-{2-[4-(4-Fluorobenzoyl)-1-piperidinyl]ethyl}-2,4-dioxo-3,4-dihydro-1(2H)-quinazolinyl]-N,N-dimethylacetamide hydrochloride,
N-{2-[4-(3,4-Dichlorobenzoyl)-1-piperazinyl]ethyl}-3-methoxybenzamide hydrochloride,
3,4-Dichloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}benzamide,
4-Chloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide hydrochloride,
N~7~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-9-methyl-9H-purin-6-amine,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzothiazol-2-amine,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzoxazol-2-amine,
6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrazinamine,
6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-pyridazinamine,
6-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione,
N-{1-[4-(3,4-Dichlorophenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-4-methyl-benzamide, hydrochloride salt,
N-{1-[4-(3,4-Dichloro-phenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-3-methoxy-benzamide, hydrochloride salt,
N-{2-[4-(3,4-Dichloroanilino)-1-piperidinyl]ethyl}-3-methoxybenzamide dihydrochloride,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N-(3-methoxybenzyl)amine dihydrochloride,
3-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methoxy-2,4(1H,3H)-quinazolinedione,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-nitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methylbenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-dinitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-iodobenzamide,
4-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
4-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methylbenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-nitrobenzamide,
3-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
3,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide,
2,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methylbenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-iodobenzamide,
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-nitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-5-(trifluoromethyl)benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide,
3,5-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)benzamide,
3-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
2-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide,
3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
2-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-difluorobenzamide,
2,3-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-naphthamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-6-(trifluoromethyl)benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-difluorobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-thiophenecarboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide,
Methyl 4-({2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}amino)-4-oxobutanoate,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclobutanecarboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxyacetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclohexanecarboxamide,

(E)-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide,

2-Chloro-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}nicotinamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclopentanecarboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenoxyacetamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide,

4-(tert-Butyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methylbenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-methylbenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-cyanobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-furamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-naphthamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(2,3-dihydro-1,4-benzodioxin-2-yl)-1,3-thiazole-4-carboxamide,

N~2~-Cyclopropyl-N~4~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine,

2-{[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl]amino}-1-ethanol,

2-[[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~-phenyl-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)-4-pyrimidinamine,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2,4-pyrimidinediamine,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine,

2-Chloro-N~4~-cyclopropyl-N~6~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenyl-2-pyrimidinamine,

N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(propylsulfanyl)-2-pyrimidinamine,

N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~-phenyl-2,4-pyrimidinediamine,

N~4~-Cyclopropyl-N~2~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}[1,8]naphthyridin-2-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-pyridinyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-dimethoxy-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-furyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-purin-6-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methylthieno[2,3-d]pyrimidin-4-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methylthieno[3,2-d]pyrimidin-4-amine,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-thiophenecarboxamide,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}cyclohexanecarboxamide,
(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-phenoxyacetamide,
(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(4-nitrophenyl)-2-propenamide,
2-(1-Adamantyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}acetamide,
(4-Chlorophenyl)(1-{2-[(2-fluoro-4,5-dimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3,4,5-trimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl){1-[2-(isobutylamino)ethyl]-4-piperidinyl}methanone,
4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-ethylhexanenitrile,
(4-Chlorophenyl)(1-{2-[(7-hydroxy-3,7-dimethyloctyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)[1-(2-{[(6-nitro-1,3-benzodioxol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,
[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,
(4-Chlorophenyl)[1-(2-{[3-nitro-4-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl)[1-(2-{[(E)-3-(4-nitrophenyl)-2-propenyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl){1-[2-({[5-(3-nitrophenyl)-2-furyl]methyl}amino)ethyl]-4-piperidinyl}methanone,
(4-Chlorophenyl)[1-(2-{[5-nitro-2-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone,
6-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-2-(methylsulfanyl)nicotinonitrile,

{1-[2-({[5-Chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazol-4-yl]methyl}amino)ethyl]-4-piperidinyl}(4-chlorophenyl)methanone,

(4-Chlorophenyl)[1-(2-{[3-(methylsulfanyl)butyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(4-phenyl-4-piperidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(1-phenyl-1H-pyrazol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,

Ethyl 3-[({2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}-amino)methyl]cyclohexanecarboxylate,

N-{4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]phenyl}acetamide,

(4-Chlorophenyl)(1-{2-[(2,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(2,6-dichlorobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(2-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)[1-(2-{[(3-methyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-methoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,

3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4H-chromen-4-one,

[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,

(4-Chlorophenyl)[1-(2-{[(2,6-dichloro-4-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(2-phenyl-1H-imidazol-4-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(5-ethyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(2-chloro-3-quinolinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(6-methyl-2-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)(1-{2-[(3-quinolinylmethyl)amino]ethyl}-4-piperidinyl)methanone,
4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-1,5-dimethyl-2-phenyl-1,2-dihydro-3H-pyrazol-3-one,
(4-Chlorophenyl)(1-{2-[(4-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(1-{2-[(2-Chloro-6-fluorobenzyl)amino]ethyl}-4-piperidinyl)(4-chlorophenyl)methanone,
[1-(2-{[(4-Bromo-1H-pyrazol-3-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,
3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-6,7-dimethyl-4H-chromen-4-one,
2-{2-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-nitrophenoxy}acetic acid,
(4-Chlorophenyl)[1-(2-{[(1-methyl-1H-benzimidazol-2-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl)[1-(2-{[(2,4-dimethoxy-5-pyrimidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(methylamino)benzamide,
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxy-4-methylbenzamide,
3-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzodioxole-5-carboxamide,
4-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluoro-4-methoxybenzamide,
5-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methyl-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,5-dimethyl-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-ethoxy-1-benzofuran-2-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-1-benzofuran-2-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methoxy-1-benzofuran-2-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-fluorophenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methoxyphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methylphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methylphenyl)acetamide,

2-(3-Bromophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,

2-(2-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,

2-(4-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(trifluoromethyl)phenyl]acetamide,

2-(3-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dimethoxyphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methoxyphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dichlorophenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-fluoro-4-methoxyphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-ethoxyphenyl)acetamide,

2-(1,3-Benzodioxol-5-yl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[4-(dimethylamino)phenyl]acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methylphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-difluorophenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methoxyphenyl)acetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenylbutanamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenylpropanamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(3-methoxyphenyl)propanamide,

2-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide,

2-(Acetylamino)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-pyridinyl)-1,3-thiazole-4-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-dimethyl-1,3-thiazole-5-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,5-dimethyl-1,3-oxazole-4-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-imidazole-4-carboxamide,

N-{2-[4-(3,4-Chlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide, hydrochloride salt,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2,6-dimethoxy-4-pyrimidinamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-6-(trifluoromethyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-methyl-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(ethylsulfanyl)-6-methyl-4-pyrimidinamine,

N~2~-Cyclopropyl-N~4~-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-methyl-4-pyrimidinamine, and

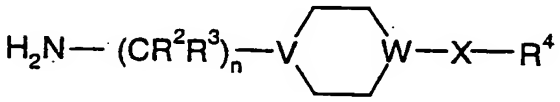
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-N~2~-phenyl-2,4-pyrimidinediamine.

The present invention further provides a process for the preparation of a compound of formula (I) which comprises

(i) when T represents a group C(O)NH, reacting a compound of general formula

$$R^1\text{-}(Q)_m\text{-}COL^1 \quad (II)$$

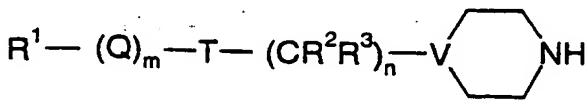
wherein L^1 represents a leaving group (e.g. a hydroxyl or halide, such as chloride, group) and R^1, m and Q are as defined in formula (I), with a compound of general formula



(III)

or an acid addition salt thereof (e.g. trifluoroacetate) wherein n, R^2, R^3, V, W, X and R^4 are as defined in formula (I); or

(ii) when T represents a group C(O)NH and W represents a nitrogen atom, reacting a compound of general formula



(IV)

wherein R^1, m, Q, T, n, R^2, R^3 and V are as defined in formula (I), with a compound of general formula

$$L^2\text{-}X\text{-}R^4 \quad (V)$$

wherein L^2 represents a leaving group (e.g. a halogen atom) and X and R^4 are as defined in formula (I); or

(iii) when T represents a group NH and m is 0, reacting a compound of general formula

$$R^1 - L^3 \quad (VI)$$

wherein L^3 represents a leaving group (e.g. a halogen atom) and R^1 is as defined in formula (I), with a compound of formula (III) as defined in (i) above; or

(iv) when T represents a group NH, m is 1 and Q represents C_1-C_4 alkylene, reacting a compound of general formula

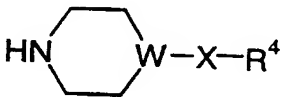
$$R^1\text{-}(CH_2)_p\text{-}CHO \quad (VII)$$

wherein p is 0, 1, 2 or 3 and R^1 is as defined in formula (I), with a compound of formula (III) as defined in (i) above; or

(v) when T represents a bond and m is 0, reacting a compound of general formula

$$R^1\text{-}(CR^2R^3)_n\text{-}L^4 \quad (VIII)$$

wherein L^4 represents a leaving group such as a halogen atom (e.g. chlorine) and n, R^1, R^2 and R^3 are as defined in formula (I), with a compound of general formula



(IX)

wherein W, X and R^4 are as defined in formula (I);

and optionally after (i), (ii), (iii), (iv) or (v) converting the compound of formula (I) to a further compound of formula (I) and/or forming a pharmaceutically acceptable salt or solvate of the compound of formula (I).

The processes of the invention may conveniently be carried out in a solvent, e.g. an organic solvent such as dimethylformamide or dichloromethane at a temperature of, for example, 15°C or above such as a temperature in the range from 20 to 100°C.

2-{[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl]amino}-1-ethanol,

2-[[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-4-pyrimidinamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2,4-pyrimidinediamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-phenyl-2-pyrimidinamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-fluoro-2,4-pyrimidinediamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(trifluoromethyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(propylsulfanyl)-2-pyrimidinamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~-phenyl-2,4-pyrimidinediamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,6-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}[1,8]naphthyridin-2-amine,

2-{[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl]amino}-1-ethanol,

2-[[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl](methyl)amino]-1-ethanol,

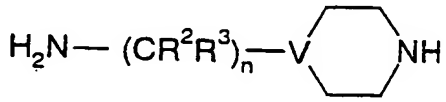
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-pyridinyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-thienyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4,6-dimethoxy-2-pyrimidinamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-furyl)-2-pyrimidinamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(2-thienyl)-2-pyrimidinamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1H-purin-6-amine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methylthieno[2,3-d]pyrimidin-4-amine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-7-methylthieno[3,2-d]pyrimidin-4-amine,
N~7~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-9-methyl-9H-purin-6-amine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,
5-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-pyridinamine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzothiazol-2-amine,
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzoxazol-2-amine,
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrazinamine,
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-3-pyridazinamine,
6-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,6-dimethoxy-4-pyrimidinamine,
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine,

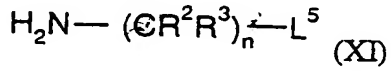
Compounds of formula (III) in which W represents a nitrogen atom may be prepared by reacting a compound of general formula



(X)

in which n, R^2, R^3 and V are as defined in formula (I) with a compound of formula (V) as defined above.

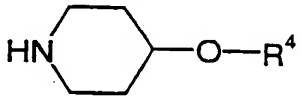
Compounds of formula (X) can be prepared by reacting piperazine with a compound of general formula



(XI)

wherein L^5 represents a halogen atom such as a bromine atom and n, R^2 and R^3 are as defined in formula (I).

Compounds of formula (III) in which W represents a group CH and X represents an oxygen atom may be prepared by reacting a compound of general formula

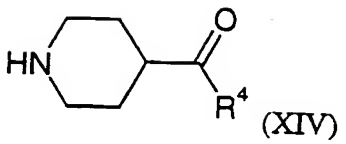


(XII)

in which R^4 is as defined in formula (I), with a compound of formula (XI).

Compounds of formula (XII) may be prepared by reacting 4-piperidinol with a compound of general formula (XIII), R^4-OH, wherein R^4 is as defined in formula (I), in the presence of a coupling agent such as diethyl azodicarboxylate and triphenylphosphine and in a solvent such as benzene or tetrahydrofuran at a temperature typically in the range from 20 to 30 °C.

Compounds of formula (III) in which W represents a group CH and X represents a group C(O) may be prepared by reacting a compound of general formula

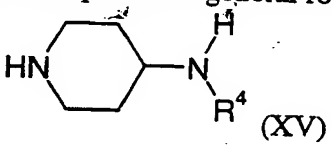


(XIV)

wherein R^4 is as defined in formula (I), with a compound of formula (XI).

Compounds of formula (III) in which W represents a group CH and X represents a group CH(OH) may be prepared by reducing/hydrogenating a corresponding compound of formula (III) in which X represents C(O) using techniques known in the art.

Compounds of formula (III) in which W represents a group CH and X represents a group NH may be prepared by reacting a compound of general formula



(XV)

in which R^4 is as defined in formula (I), with a compound of formula (XI).

Compounds of formula (XV) may be prepared by reacting 4-piperidone with a compound of general formula (XVI), R^4-NH_2, wherein R^4 is as defined in formula (I), in the presence of a reducing agent such as sodium cyanoborohydride or sodium borohydride and in a solvent such as methanol or benzene at a temperature typically in the range from 20 to 90 °C.

Compounds of formula (III) in which W represents a group CH and X represents a group N(C_1-C_6 alkyl) may be prepared by alkylating a corresponding compound of formula (III) in which X represents a group NH, using techniques conventional in the art.

Compounds of formula (IV) may be prepared by reacting a compound of formula (II) with a compound of formula (X).

Compounds of formulae II, V, VI, VII, VIII, IX, XI, XIII, XIV and XVI are either commercially available, are well known in the literature or may be prepared easily using known techniques.

Compounds of formula (I) can be converted into further compounds of formula (I) using standard procedures. For example, compounds of formula (I) in which R^1 represents an alkoxy-substituted phenyl group can be converted to compounds of formula (I) in which R^1 represents a hydroxy-substituted phenyl group by reaction with boron tribromide in a solvent such as dichloromethane. Further, compounds of formula (I) in which X represents C(O) can be converted to compounds of formula (I) in which X represents CH(OH) by reaction with triethylsilane and trifluoroacetic acid in a solvent such as dichloromethane.

It will be appreciated by those skilled in the art that in the processes of the present invention certain functional groups such as hydroxyl or amino groups in the starting reagents or intermediate compounds may need to be protected by protecting groups. Thus, the preparation of the compounds of formula (I) may involve, at an appropriate stage, the removal of one or more protecting groups.

The protection and deprotection of functional groups is described in 'Protective Groups in Organic Chemistry', edited by J.W.F. McOmie, Plenum Press (1973) and 'Protective Groups in Organic Synthesis', 2nd edition, T.W. Greene and P.G.M. Wuts, Wiley-Interscience (1991).

The compounds of formula (I) above may be converted to a pharmaceutically acceptable salt or solvate thereof, preferably an acid addition salt such as a hydrochloride, hydrobromide, phosphate, acetate, fumarate, maleate, tartrate, citrate, oxalate, methanesulphonate or *p*-toluenesulphonate.

Certain compounds of formula (I) are capable of existing in stereoisomeric forms. It will be understood that the invention encompasses the use of all geometric and optical isomers

of the compounds of formula (I) and mixtures thereof including racemates. The use of tautomers and mixtures thereof also form an aspect of the present invention.

The compounds of formula (I) have activity as pharmaceuticals, in particular as modulators of chemokine receptor (especially CCR1 and/or CCR3) activity, and may be used in the treatment of autoimmune, inflammatory, proliferative and hyperproliferative diseases and immunologically-mediated diseases including rejection of transplanted organs or tissues and Acquired Immunodeficiency Syndrome (AIDS).

Examples of these conditions are:

(1) **(the respiratory tract)** obstructive airways diseases including chronic obstructive pulmonary disease (COPD) such as irreversible COPD; asthma, such as bronchial, allergic, intrinsic, extrinsic and dust asthma, particularly chronic or inveterate asthma (e.g. late asthma and airways hyper-responsiveness); bronchitis; acute, allergic, atrophic rhinitis and chronic rhinitis including rhinitis caseosa, hypertrophic rhinitis, rhinitis purulenta, rhinitis sicca and rhinitis medicamentosa; membranous rhinitis including croupous, fibrinous and pseudomembranous rhinitis and scrofoulous rhinitis; seasonal rhinitis including rhinitis nervosa (hay fever) and vasomotor rhinitis; sarcoidosis, farmer's lung and related diseases, fibroid lung and idiopathic interstitial pneumonia;

(2) **(bone and joints)** rheumatoid arthritis, seronegative spondyloarthropathies (including ankylosing spondylitis, psoriatic arthritis and Reiter's disease), Behcet's disease, Sjogren's syndrome and systemic sclerosis;

(3) **(skin)** psoriasis, atopical dermatitis, contact dermatitis and other eczmatous dermitides, seborrhoetic dermatitis, Lichen planus, Pemphigus, bullous Pemphigus, Epidermolysis bullosa, urticaria, angiodermas, vasculitides, erythemas, cutaneous eosinophilias, uveitis, Alopecia areata and vernal conjunctivitis;

(4) **(gastrointestinal tract)** Coeliac disease, proctitis, eosinopilic gastro-enteritis, mastocytosis, Crohn's disease, ulcerative colitis, food-related allergies which have effects remote from the gut, e.g., migraine, rhinitis and eczema;

(5) **(other tissues and systemic disease)** multiple sclerosis, atherosclerosis, Acquired Immunodeficiency Syndrome (AIDS), lupus erythematosus, systemic lupus, erythematosus, Hashimoto's thyroiditis, myasthenia gravis, type I diabetes, nephrotic syndrome, eosinophilia fascitis, hyper IgE syndrome, lepromatous leprosy, sezary syndrome and idiopathic thrombocytopenia pupura; and

(6) **(allograft rejection)** acute and chronic following, for example, transplantation of kidney, heart, liver, lung, bone marrow, skin and cornea; and chronic graft versus host disease.

Thus, the present invention provides a compound of formula (I), or a pharmaceutically-acceptable salt or solvate thereof, as hereinbefore defined for use in therapy.

In a further aspect, the present invention provides the use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined in the manufacture of a medicament for use in therapy.

In the context of the present specification, the term "therapy" also includes "prophylaxis" unless there are specific indications to the contrary. The terms "therapeutic" and "therapeutically" should be construed accordingly.

The invention also provides a method of treating an inflammatory disease in a patient suffering from, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined.

For the above-mentioned therapeutic uses the dosage administered will, of course, vary with the compound employed, the mode of administration, the treatment desired and the disorder indicated.

The compounds of formula (I) and pharmaceutically acceptable salts and solvates thereof may be used on their own but will generally be administered in the form of a pharmaceutical composition in which the formula (I) compound/salt/solvate (active ingredient) is in association with a pharmaceutically acceptable adjuvant, diluent or carrier. Depending on the mode of administration, the pharmaceutical composition will preferably comprise from 0.05 to 99 %w (per cent by weight), more preferably from 0.05 to 80 %w, still more preferably from 0.10 to 70 %w, and even more preferably from 0.10 to 50 %w, of active ingredient, all percentages by weight being based on total composition.

The present invention also provides a pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined, in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

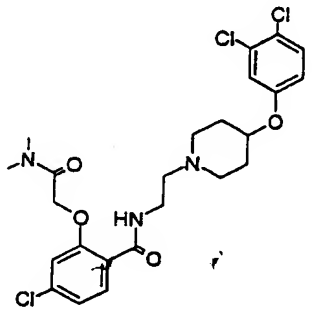
The invention further provides a process for the preparation of a pharmaceutical composition of the invention which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined, with a pharmaceutically acceptable adjuvant, diluent or carrier.

The pharmaceutical compositions may be administered topically (e.g. to the lung and/or airways or to the skin) in the form of solutions, suspensions, heptafluoroalkane aerosols and dry powder formulations; or systemically, e.g. by oral administration in the form of tablets, capsules, syrups, powders or granules, or by parenteral administration in the form of solutions or suspensions, or by subcutaneous administration or by rectal administration in the form of suppositories or transdermally.

The invention will now be further explained by reference to the following illustrative examples.

Example 1

4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide



(i) tert-Butyl 4-(3,4-dichlorophenoxy)-1-piperidinecarboxylate

Diethyl azodicarboxylate (12.6ml) was added to a solution of triphenylphosphine (20.8g) in tetrahydrofuran (300ml) at 0° C. After 15 minutes 3,4-dichlorophenol (12.9g) was added, after a further 10 minutes tert-butyl 4-hydroxy-1-piperidinecarboxylate (14.5g) in tetrahydrofuran (100ml) was added dropwise over 0.5 hour. The solution was stirred at room temperature for 5 hours and concentrated to a small volume. The residue was partitioned between ether and brine. The organic phase was separated, dried and evaporated to a gum. Purification by chromatography (ethyl acetate : isohexane 95:5) gave the sub-titled product as an oil (20g).

MS: APCI(+ve): 246 (M-BOC+2H)

(ii) 4-(3,4-Dichlorophenoxy)piperidine

The product from step (i) was dissolved in dichloromethane (200ml) and trifluoroacetic acid (100ml) was added. After 18 hours at room temperature the solution was evaporated and the resultant gum triturated under ether to give the sub-titled product as a solid (16.2g).

(iii) tert-Butyl 2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethylcarbamate

The product from step (ii) (6.55g) was dissolved in DMF (50ml) and triethylamine (7.9ml) was added. tert-Butyl 2-bromoethylcarbamate (4.3 g) in DMF (5ml) was added and the solution stirred at room temperature for 3 days. Ethyl acetate and water were added, the organic phase separated, dried and evaporated to a gum. Purification by chromatography (dichloromethane : methanol 95:5) gave the sub-titled product as a gum (5.7g).

MS: APCI(+ve): 389(M+H)

(iv) 2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethylamine trifluoroacetate

The product from step (iii) was dissolved in dichloromethane (200ml) and trifluoroacetic acid (100ml) added. After 18hrs at room temperature the solvent was evaporated and the resultant gum triturated under ether to give the sub-titled product as a solid (5.7g).

MS: APCI(+ve): 290(M+H)

(v) 2-(Dimethylamino)-2-oxoethyl 4-chloro-2-[2-(dimethylamino)-2-oxoethoxy]benzoate

A mixture of 4-chloro-2-hydroxybenzoic acid (5g), Cs_2CO_3 (17.5g) and 2-chloro-N,N-dimethylacetamide (6.6g) was stirred and heated at 70 °C for 3 hours. Water and ethyl acetate were added, the organic phase separated, dried and concentrated to a gum which was purified by chromatography (ethyl acetate : methanol, 9:1) to give the sub-titled product as a solid (8.0g).

MS: APCI(+ve) 343(M+H)

Melting point: 140-141 °C

(vi) 4-Chloro-2-[2-(dimethylamino)-2-oxoethoxy]benzoic acid

The product from step (v) (1.0g) was dissolved in a 2 : 1 water : methanol mixture (15ml) and $LiOH.H_2O$ added. After 2 hours 2M aqueous HCl solution and ethyl acetate were

added, the organic phase separated, dried and concentrated to give the sub-titled product as a solid (1.2g).

MS: APCI(+ve) 258(M+H)

Melting point: 141-142 °C

(vii) 4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide

The product from step (vi) (0.3g) and N,N-carbonyldiimidazole (0.19g) were dissolved in DMF (20ml) and the solution stirred at room temperature for 1 hour. The product from step (iv) (0.42g) and triethylamine (0.32ml) were added. After 20 hours water and ether were added, the organic phase separated, dried and concentrated to a gum which was purified by chromatography (dichloromethane : methanol, 93:7) to give the titled product as a solid (0.38g).

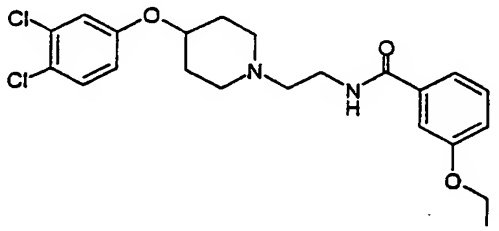
MS: ESI 528.12 (M+H)

^{1}H NMR: δ(DMSO) 9.17 (t, 1H), 7.88 (d, 1H), 7.48 (d, 1H), 7.38 (d, 1H), 7.24 (d, 1H), 7.13 (dd, 1H), 6.99 (dd, 1H), 5.11 (s, 2H), 4.32 (m, 1H), 3.42 (m, 2H), 2.99 (s, 3H), 2.88 (s, 3H), 2.73 (m, 2H), 2.50 (m, 2H), 2.30 (m, 2H), 1.90 (m, 2H), 1.59 (m, 2H).

Melting point: 139-40 °C

Example 2

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-ethoxybenzamide hydrochloride



The product of Example 1 step (iv) (0.4g) was dissolved in DMF (10ml) , PyBrop (0.541g), 3-ethoxybenzoic acid (0.167g) and N,N-di-isopropylethylamine (0.5g) were added. After

18 hours at room temperature chloroform and aqueous $NaHCO_3$ solution were added. The organic phase was separated, dried and concentrated to leave a gum which was purified by chromatography (ethyl acetate : methanol 97:3) to give an oil. Addition of 1.0M ethereal hydrogen chloride solution gave the titled product as a solid (0.14g).

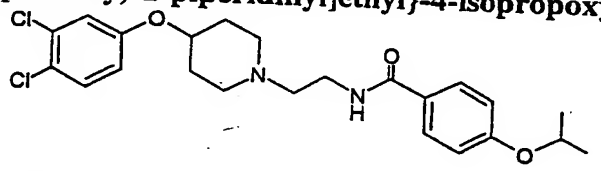
MS: ESI 437.14 (M+H)

^{1}H NMR: δ(DMSO) 8.87 (bs, 1H), 7.50 (m, 3H), 7.40 (m, 2H), 7.06 (m, 2H), 4.83 / 4.62 (m, 1H), 4.08 (q, 2H), 3.67 (m, 3H), 3.47 (m, 1H), 3.17 (m, 3H), 2.20 (m, 2H), 2.03 (m, 2H), 1.34 (t, 3H)

Melting point: 191-193 °C

Example 3

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-isopropoxybenzamide



Prepared by the same method as Example 2 using 4-isopropoxybenzoic acid without the addition of 1.0 M ethereal hydrogen chloride solution to give the titled product as a solid (0.12g).

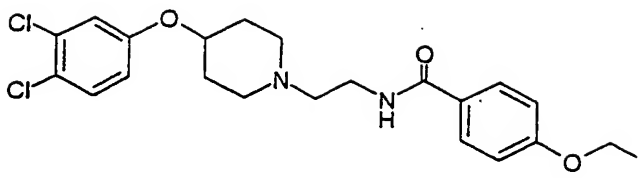
MS: ESI 451.14 (M+H)

^{1}H NMR: δ(DMSO) 8.22 (t, 1H), 7.8 (m, 2H), 7.49 (d, 1H), 7.25 (d, 1H), 7.00 (m, 3H), 4.7 (m, 1H), 4.45 (m, 1H), 3.36 (m, 2H), 2.73 (m, 2H), 2.48 (m, 2H), 2.29 (m, 2H), 1.91 (m, 2H), 1.60 (m, 2H), 1.28 (s, 3H), 1.27 (s, 3H)

Melting point: 113-15 °C

Example 4

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-ethoxybenzamide



Prepared by the same method as Example 2 using 4-ethoxybenzoic acid without the addition of 1.0 M ethereal hydrogen chloride solution to give the titled product as a solid (0.1g).

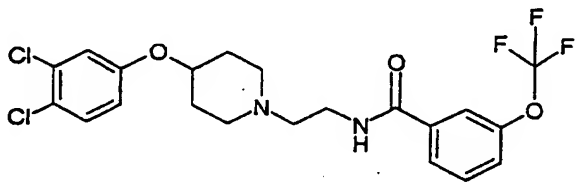
MS: ESI 437.14 (M+H)

^{1}H NMR: δ(DMSO) 8.22 (t, 1H), 7.79 (d, 2H), 7.49 (d, 1H), 7.25 (d, 1H), 7.00 (m, 3H), 4.5 (m, 1H), 4.07 (q, 2H), 3.37 (q, 2H), 2.73 (m, 2H), 2.47 (m, 2H), 2.30 (m, 2H), 1.91 (m, 2H), 1.60 (m, 2H), 1.34 (t, 3H)

Melting point: 118-20 °C

Example 5

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide hydrochloride



Prepared by the same method as Example 2 using 3-trifluoromethoxybenzoic acid to give the titled product as a solid (0.12g).

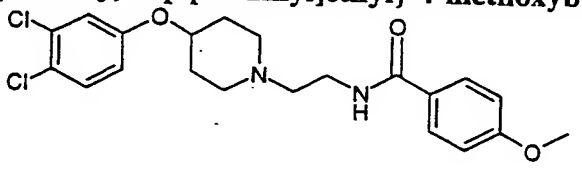
MS: ESI 477.09 (M+H)

^{1}H NMR: δ(DMSO) 10.42 (bs, 1H), 9.11 (bm, 1H), 8.0 (d, 1H), 7.88 (s, 1H), 7.6 (m, 3H), 7.37 (m, 1H), 7.06 (m, 1H), 4.70 (m, 1H), 3.71 (m, 3H), 3.48 (d, 1H), 3.20 (m, 4H), 2.2 (m, 4H)

Melting point: 180-82 °C

Example 6

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide



Prepared by the same method as Example 2 using 4-methoxybenzoic without the addition of 1.0 M ethereal hydrogen chloride solution to give the titled product as a solid (0.11g).

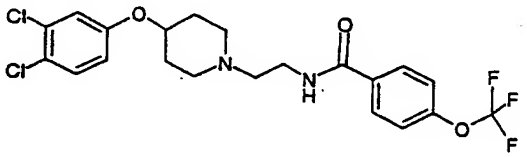
MS: ESI 423.12 (M+H)

^{1}H NMR: δ(DMSO) 8.42 (t, 1H), 7.81 (m, 2H), 7.49 (d, 1H), 7.25 (d, 1H), 6.98 (s, 3H), 4.4 (m,1H), 3.8 (s, 3H), 3.35 (q, 2H), 2.73 (m, 2H), 2.47 (m, 2H), 2.30 (m, 2H), 1.91 (m, 2H), 1.60 (m, 2H)

Melting point: 110-12 °C

Example 7

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethoxy)benzamide hydrochloride



Prepared by the same method as Example 2 using 4-trifluoromethoxybenzoic acid to give the titled product as a solid (0.19g).

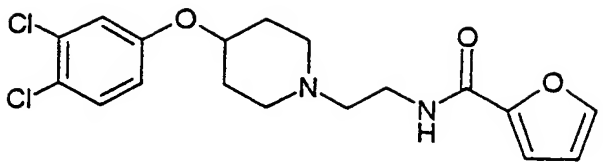
MS: ESI 477 (M+H)

^{1}H NMR: δ(DMSO) 10.5 (bs, 1H), 9.06 (m, 1H), 8.07 (dd, 2H), 7.55 (t, 1H), 7.49 (d, 2H), 7.36 (dd, 1H), 7.10-7.02 (m, 1H), 4.72 (m, 1H), 3.70 (m, 3H), 3.47 (d, 1H), 3.14 (m, 2H), 2.25 (m, 2H), 2.02 (m, 2H)

Melting point: 184-187 °C

Example 8

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide hydrochloride



Prepared by the same method as Example 2 using furan-2-carboxylic acid to give the titled product as a solid (0.14g).

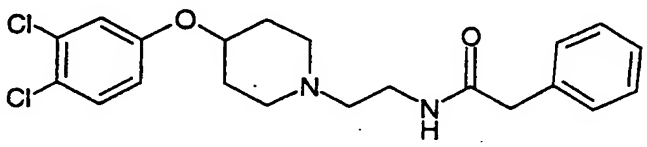
MS: ESI 383.09 (M+H)

^{1}H NMR: δ(DMSO) 10.43 (bm, 1H), 8.76 (t, 1H), 7.87 (s, 1H), 7.55 (t, 1H), 7.36 (dd, 1H), 7.21 (d, 1H), 7.06 (m, 1H), 6.64 (dd, 1H), 4.83-4.61 (m, 1H), 3.65 (m, 3H), 3.45 (d, 1H), 3.08 (m, 4H), 2.1 (m, 4H)

Melting point: 225-28 °C

Example 9

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide hydrochloride



Prepared by the same method as Example 2 using phenylacetic acid to give the titled product as a solid (0.12g).

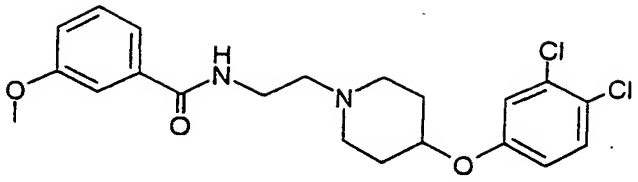
MS: ESI 407 (M+H)

^{1}H NMR: δ(DMSO) 10.28 (bm, 1H), 8.46 (bm, 1H), 7.56 (t, 1H), 7.3 (m, 6H), 7.10 (m, 1H), 4.81/4.58 (m, 1H), 3.58 (d, 1H), 3.46 (m, 4H), 3.10 (m, 4H), 2.15 (m, 5H)

Melting point: 135-38 °C

Example 10

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride



The product of Example 1 step (iv) (2.0g) was dissolved in dichloromethane (490ml), triethylamine (1.85ml) and 3-methoxybenzoyl chloride (0.66g) were added. After 72 hours at room temperature, water was added, the organic phase separated, dried and concentrated to a gum. The product was dissolved in dichloromethane and treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.88g).

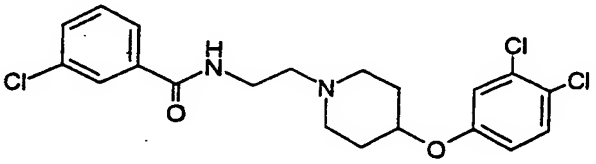
MS: ESI 423.12 (M+H)

^{1}H NMR: δ(DMSO) 10.6-10.5 (m, 1H), 9.92 (s, 1H), 7.54 (m, 3H), 7.38 (m, 2H), 7.08 (m, 2H), 4.84/4.62 (m 1H), 3.82 (s, 3H), 3.45 (m, 8H), 2.27 (m, 4H).

Melting point: 72-73 °C

Example 11

3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide hydrochloride



The product of Example 1 step (iv) (0.15g) was dissolved in DMF (3ml), N,N-di-isopropylethylamine (0.3ml) and 3-chlorobenzoyl chloride (0.054ml) were added. After 2 hours at room temperature, water and ethyl acetate were added, the organic phase separated dried and concentrated. The residue was purified by chromatography (dichloromethane : methanol, 95:5) to give an oil which was dissolved in ether and 1.0M ethereal hydrogen chloride solution added to give the titled product as a solid (0.12g).

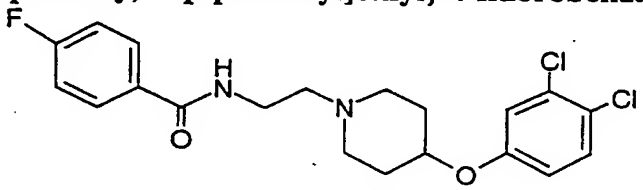
MS: ESI 427.07 (M+H)

^{1}H NMR: δ(DMSO) 8.42 (t, 1H), 7.94-7.84 (m, 2H), 7.49 (d, 1H), 7.29 (m, 3H), 6.98 (dd, 1H), 4.44 (m, 1H), 3.36 (m, 2H), 2.74 (m, 2H), 2.48 (m, 2H), 2.29 (bt, 2H), 1.92 (m, 2H), 1.60 (m, 2H)

Melting point: 118 °C

Example 12

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide



Prepared by the same method as Example 11 using 4-fluorobenzoyl chloride without the addition of 1.0 M ethereal hydrogen chloride solution to give the titled product as a solid (0.1g).

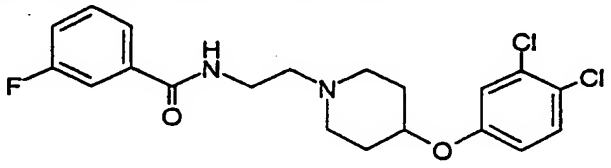
MS: ESI 411.10 (M+H)

^{1}H NMR: δ(DMSO) 10.46 (bs, 1H), 9.04 (s, 1H), 7.98 (s, 1H), 7.90 (d, 1H), 7.58 (m, 3H), 7.36 (dd, 1H), 7.05 (m, 1H), 4.84/4.60 (m, 1H), 3.69 (m, 3H), 3.48 (bd, 1H), 3.20 (m, 4H), 2.15 (m, 4H)

Melting point: 192 °C

Example 13

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide hydrochloride



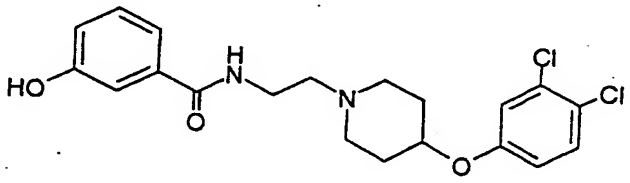
Prepared by the same method as Example 11 using 3-fluorobenzoyl chloride to give the titled product as a solid (0.09g).

MS: ESI 411.10 (M+H)

^{1}H NMR: δ(DMSO) 10.67 (bs, 1H), 9.06 (s, 1H), 7.80 (m, 2H), 7.55 (m, 2H), 7.40 (m, 2H), 7.05 (m, 1H), 4.84/4.63(m, 1H), 3.70 (m, 3H), 3.28 (m, 3H), 2.20 (m, 4H)
Melting point: 225 °C

Example 14

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-hydroxybenzamide hydrochloride

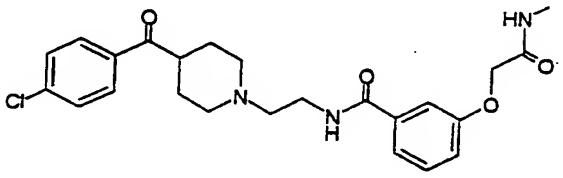


The product of Example 10 (0.15g) was dissolved in dichloromethane (10ml) and a solution of 1.0M BBr_3 in dichloromethane (4ml) added. After 16 hours at room temperature the solvent was removed by evaporation, methanol was added and the solution concentrated. The residue was dissolved in 2M aqueous HCl solution, concentrated to dryness and the residue triturated under ether to give the titled product as a solid (0.1g).

MS: ESI 409.10 (M+H)
^{1}H NMR: δ(DMSO) 9.98-9.4 (bs, 2H), 8.71 (t, 1H), 7.6 (dd, 1H), 7.4-7.2 (m, 4H), 7.05 (m, 1H), 6.95 (dd, 1H), 4.65 (m, 1H), 3.40 (m, 8H), 2.0 (m, 4H)
Melting point: 83-4 °C

Example 15

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-[2-(methylamino)-2-oxoethoxy]benzamide hydrochloride



(i) **[1-(2-Aminoethyl)-4-piperidinyl](4-chlorophenyl)methanone trifluoroacetate**

To a solution of (4-chlorophenyl)(4-piperidinyl)methanone hydrochloride (2.5g) and tert-butyl 2-bromoethylcarbamate (2.1g) in DMF was added triethylamine (2.9g), after 72 hours at room temperature water and ether were added. The organic phase was separated, dried and concentrated. The residue was dissolved in dichloromethane (40 ml), trifluoroacetic acid (10ml) added and the solution left for 20 hours. Evaporation of the solvent gave a sticky solid which was triturated under ether to give the sub-titled product as a solid (2.5g).

(ii) N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride

The product of step (i) (2.5g) was dissolved in dichloromethane (20ml), triethylamine (0.75ml) and 3-methoxybenzoyl chloride (0.276g) were added. After 16 hours, water was added, the organic phase separated, dried and concentrated to a gum. Purification by chromatography (ethyl acetate) gave a gum, which was treated with 1.OM ethereal hydrogen chloride solution to give the sub-titled product as a solid (0.3g).

MS: ESI 401.16 (M+H)

^{1}H NMR: δ(DMSO) 10.3 (bm, 1H), 8.95 (t, 1H), 8.0 (m, 2H), 7.6 (m, 2H), 7.5 (m, 2H), 7.4 (t, 1H), 7.05 (m, 1H), 3.8 (s, 3H), 3.68 (m, 4H), 3.28 (m, 5H), 2.0 (m, 4H).

Melting point: 196-7 °C

(iii) N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-hydroxybenzamide hydrochloride

Prepared by the method of Example 14 using the product of step (ii) above (0.24g) to give the sub-titled product as a solid (0.20g).

MS: ESI 387.14 (M+H)

^{1}H NMR: δ(DMSO) 8.62 (t, 1H), 8.05 (dd, 2H), 7.6 (dd, 2H), 7.25 (m, 3H), 6.95 (m, 1H), 4.26 (m, 9H), 2.0 (m, 4H)

Melting point: 90-91 °C

(iv) N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-[2-(methylamino)-2-oxoethoxy]benzamide hydrochloride

The product of step (iii) above (0.10g) was dissolved in DMF (3ml), Cs_2CO_2 (0.23g) and 2-chloro-N-methylacetamide (0.26g) were added and the mixture heated at 80° C for 16 hours. The mixture was cooled to room temperature, water and ethyl acetate were added and the organic phase separated. Evaporation of the solvent gave a gum which was treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.05g).

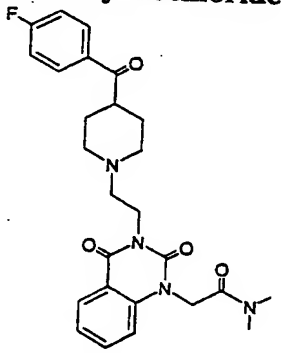
MS: ESI 458.18 (M+H)

^{1}H NMR: δ(DMSO) 10.6-10.2 (bm, 1H), 8.95 (bm, 1H), 8.1 (m, 2H), 7.55 (m, 8H), 7.14 (bd, 1H), 4.54 (s, 2H), 4.0 (m, 1H), 3.4 (m, 8H), 2.65 (d, 3H), 2.0 (m, 4H)

Melting point: 69-70 °C

Example 16

2-[3-{2-[4-(4-Fluorobenzoyl)-1-piperidinyl]ethyl}-2,4-dioxo-3,4-dihydro-1(2H)-quinazolinyl]-N,N-dimethylacetamide hydrochloride



3-{2-[4-(4-Fluorobenzoyl)-1-piperidinyl]ethyl}-2,4(1H,3H)-quinazolinedione was dissolved in DMF (5ml) and NaH (60% dispersion in mineral oil) added. After 0.5 hours, 2-chloro-N,N-dimethylacetamide was added and the solution stirred at room temperature for 16 hours. Water and ethyl actetate were added, the organic phase separated, dried and concentrated to an oil. Purification by chromatography (dichloromethane : methanol 95:5) gave an oil which was treated with 1.0 M ethereal hydrogen chloride solution to give the titled product as a solid(0.015g).

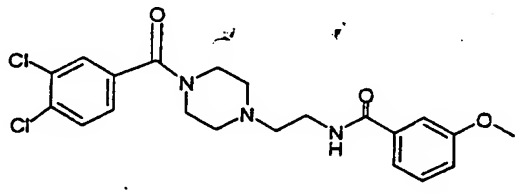
MS: ESI 481.22 (M+H)

^{1}H NMR: δ(DMSO) 8.08 (m,3H), 7.76 (t, 1H), 7.40 (t, 2H), 7.32 (m, 2H), 5.05 (s, 2H), 4.36 (m, 1H), 3.76 (m, 3H), 3.39 (m, 2H), 3.15 (s, 3H), 2.87 (s, 3H), 2.02 (m, 2H), 1.81 (m, 2H), 1.28 (m,2H)

Melting point: 245-246 °C

Example 17

N-{2-[4-(3,4-Dichlorobenzoyl)-1-piperazinyl]ethyl}-3-methoxybenzamide hydrochloride



(i) tert-Butyl 2-(1-piperazinyl)ethylcarbamate

A mixture of benzaldehyde (21g) and 1-(2-aminoethyl)piperazine (25.8g) was stirred and heated under a Dean and Stark water separator for 20 hours. The cooled solution was treated portionwise with di-tert-butyldicarbonate (48g), stirred for 72 hours and concentrated. The residue was treated with 1M aqueous $KHSO_4$ solution (220ml), stirred for 24hours, ether was added and the organic phase separated. The aqueous phase was treated with 2M NaOH solution, dichloromethane was added and the organic phase separated. The combined organic phase was washed with brine, dried and concentrated to give the sub-titled product as an oil (30g).

MS: APCI(+ve) 230(M+H)

^{1}H NMR δ ($CDCl_3$) 3.43 (t, 4H), 2.8 (t, 2H), 2.45 (m, 6H), 1.5 (s, 9H).

(ii) tert-Butyl 2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethylcarbamate

The product from step (i) above (3g) was dissolved in pyridine (12ml), 3,4-dichlorobenzoyl chloride (2.05g) was added and the mixture stirred at room temperature for 18 hours. A

solid was collected by filtration and purified by chromatography (dichloromethane : methanol : 0.880 NH_4OH, 90:9:1) to give the sub-titled product as an oil (3.59g).

MS: APCI(+ve) 364(M+H)

^1H NMR δ ($CDCl_3$) 7.33 (m, 3H), 7.04 (m, 1H), 6.76 (bs, 1H), 3.86 (s, 3H), 3.55 (q, 2H), 3.45 (t, 4H), 2.61 (t, 3H), 2.46 (t, 4H), 1.46 (s, 9H)

(iii) [4-(2-Aminoethyl)-1-piperazinyl](3,4-dichlorophenyl)methanone trifluoroacetate

The product from step (ii) above (3.3g) was dissolved in dichloromethane (50ml) and trifluoroacetic acid (10ml) added. After 16 hours at room temperature the solvent was removed to give the sub-titled product as an oil (5.9g).

MS: APCI(+ve) 264(M+H)

(iv) N-{2-[4-(3,4-Dichlorobenzoyl)-1-piperazinyl]ethyl}-3-methoxybenzamide hydrochloride

The product from step (iii) above (0.15g) was dissolved in pyridine (2ml) and 3-methoxybenzoyl chloride (0.064g) added. After 16 hours at room temperature, water and ethyl acetate were added, the organic phase separated, dried and concentrated to an oil. Purification by chromatography (dichloromethane : methanol, 95:5) gave an oil which was treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.043g).

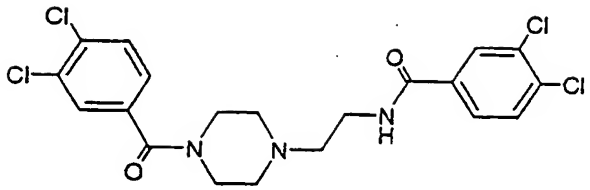
MS: ESI 436.12 (M+H)

^1H NMR: δ(DMSO) 8.8 (bt, 1H), 7.34 (m, 2H), 7.43 (m, 4H), 7.14(m, 1H), 3.82(s, 3H), 3.48 (m,12H)

Melting point: 230 °C

Example 18

3,4-Dichloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}benzamide



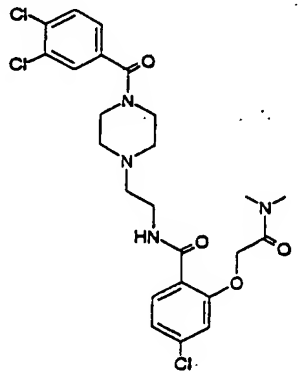
A solution of benzaldehyde (5.3g) and 1-(2-aminopiperazine) (6.45g) in toluene (100ml) was heated under a Dean and Stark water separator for 4 hours. The solution was cooled to room temperature and triethylamine (5.05g) added. A solution of 3,4-dichlorobenzoyl chloride (10.48g) in toluene (50ml) was added dropwise, the solution stirred at room temperature for 18 hours and water added. The organic phase was separated, dried and concentrated to a residue which was treated with 1N aqueous $KHSO_4$ solution (65ml). The mixture was stirred vigorously for 4 hours, ether was added, the aqueous phase separated and NaOH added. $CHCl_3$ was added, the organic phase separated, dried and concentrated to a gum. Purification by chromatography (dichloromethane : triethylamine, 95:5) gave the titled product as a foam (0.25g).

MS: ESI 474.03 (M+H)

^{1}H NMR: δ(DMSO) 8.8 (bt, 1H), 7.34 (m, 2H), 7.43 (m, 4H), 7.14 (m, 1H), 3.82 (s, 3H), 3.48 (m, 12H)

Example 19

4-Chloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide hydrochloride



The product of Example 26 step (ii) (0.3g), 3,4-dichlorobenzoyl chloride (0.1g) and triethylamine (0.15g) were dissolved in dichloromethane (15ml). After 20 hours at room

temperature water was added, the organic phase separated, dried and evaporated to give a gum. Purification by chromatography (dichloromethane : methanol, 20:1) gave a solid which was treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.1g).

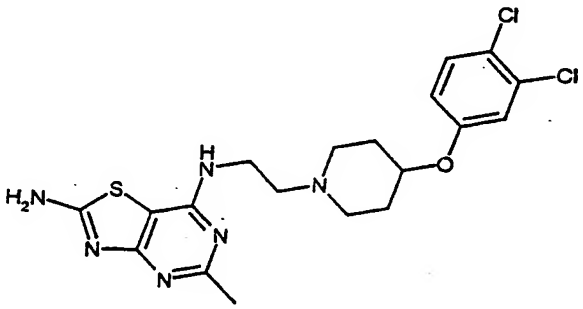
MS: ESI 541.11 (M+H)

^{1}H NMR δ(DMSO-D6) 9.54 (t, 1H), 7.91 (d, 1H), 7.74 (m, 2H), 7.43 (m, 2H), 7.18 (d, 1H), 5.12 (s, 2H), 3.2-3.8 (m, 12H), 2.99 (s, 3H), 2.88 (s,3H).

Melting point: 226-7 °C.

Example 20

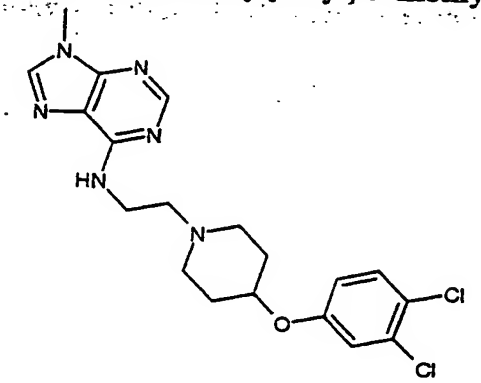
N~7~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine



MS: APCI(+ve) 453 (M+1)

Example 21

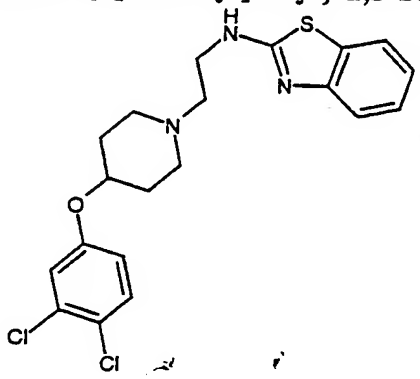
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-9-methyl-9H-purin-6-amine



MS: APCI(+ve) 421 (M+1)

Example 22

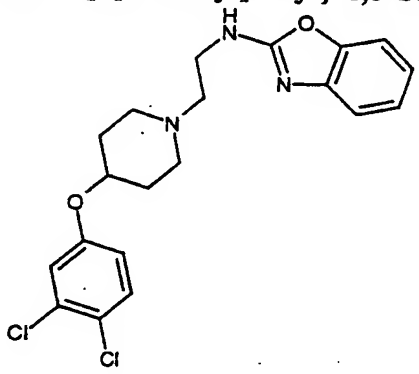
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzothiazol-2-amine



MS: APCI(+ve) 422 (M+1)

Example 23

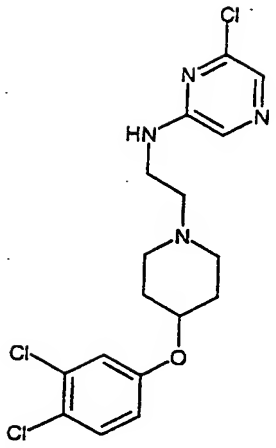
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzoxazol-2-amine



MS: APCI(+ve) 406 (M+1)

Example 24

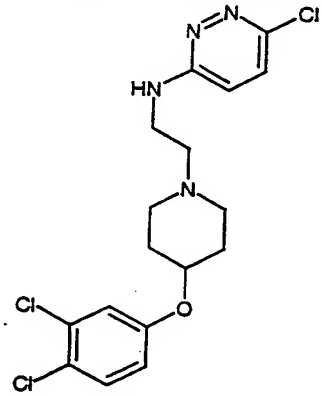
6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrazinamine



MS: APCI(+ve) 403 (M+1)

Example 25

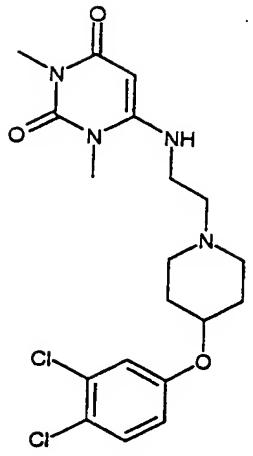
6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-pyridazinamine



MS: APCI(+ve) 403 (M+1)

Example 26

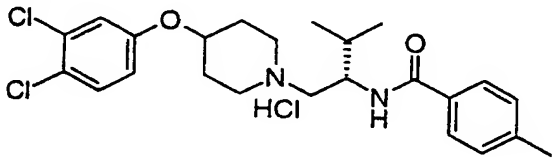
6-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione



MS: APCI(+ve) 427 (M+1)

Example 27

N-{1-[4-(3,4-Dichlorophenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-4-methyl-benzamide, hydrochloride



(i) N-{1-{4-(3,4-Dichlorophenoxy)-piperidine-1-carbonyl]-2-methyl-propyl}-acetamide

N-Boc Valine (1.13g) was dissolved in dichloromethane (5 ml) and EDC (0.99g) added, after 5 min the product according to Example 1 step (ii) (1.44g) in dichloromethane (5 ml) was added in one portion. After 3 hours at room temperature, aqueous sodium bicarbonate solution and ethyl acetate were added. The organic phase was separated and the solvent removed to give the sub-titled compound as an oil (1.57 g) which was used in the next step without further purification.

(ii) 2-amino-1-[4-(3,4-dichlorophenoxy)-piperidine-1-yl]-3-methyl-butan-1-one

The product of step (i) (1.57 g) was dissolved in dichloromethane (14 ml) and trifluoroacetic acid (4 ml) added. After 2 hours at room temperature the solvent was removed, ethyl acetate and 2N aqueous NaOH solution were added to give pH 8.0. The organic phase was separated and concentrated to give the sub-titled product as an oil (1.24 g) which was used in the next step without further purification.

(iii) 1-[4-(3,4-Dichlorophenoxy)-piperidin-1-ylmethyl]-2-methyl-propylamine

The product of step (ii) (1.12g) was dissolved in THF (10 ml) and Borane/THF complex (22.7 ml) added. The mixture was heated under reflux for 2 hours and cooled. The solvent was evaporated, the product dissolved in methanol (5ml) and 50% aqueous HCl solution added. The mixture was heated to 70 °C for 1 hour and cooled to room temperature. The solvent was removed, ethyl acetate and 2N aqueous NaOH solution were added to give pH 9.0. The organic phase was separated and the solvent evaporated to give the sub-titled compound as an oil (0.98 g) which used without further purification.

(iv) N-{1-[4-(3,4-Dichlorophenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-4-methyl-benzamide, hydrochloride

The product of step (iii) (0.2g) was dissolved in dichloromethane (5ml), triethylamine (0.126 ml) and 4-methylbenzoyl chloride (0.097 ml) were added. After 2 hours at room temperature, ethyl acetate and aqueous $NaHCO_3$ solution were added, the organic phase separated and the solvent removed to leave an oil. Purification by reverse phase HPLC (with a gradient eluent system (25% $MeCN/NH_4OAc_{aq}$ (0.1%) to 95% $MeCN//NH_4OAc_{aq}$ (0.1%)) gave a gum. Addition of 1.0M ethereal hydrogen chloride solution gave the titled product as a solid (0.104 g).

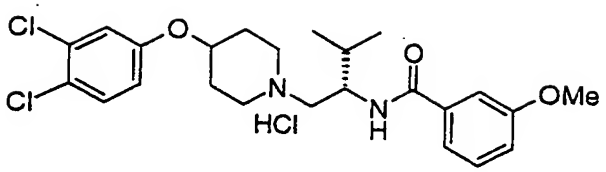
Melting point: 131-132°C

MS: ESI 450 (M+H)

^1H NMR: δ(DMSO) 8.45 (t, 1H), 7.00-7.90 (m, 7H), 4.79 (br s, 1H), 4.24-4.30 (m, 1H), 3.10-3.42 (m, 5H), 2.36 (s, 3H), 1.88-2.40 (m, 5H), 0.92 (t, 6H)

Example 28

N-{1-[4-(3,4-Dichloro-phenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-3-methoxy-benzamide, hydrochloride



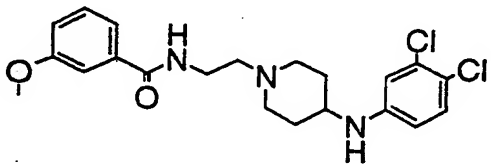
The product according to Example 27 step (iii) dissolved in dichloromethane (4 ml), triethylamine (0.090 ml) and 3-methoxybenzoyl chloride (0.077 ml) were added. After 2 hours at room temperature, $NaHCO_3$ was added, the product extracted with ethyl acetate, the combined organic extracts dried with Na_2SO_4 and concentrated. Purification with reverse phase HPLC (with a gradient eluent system (25% $MeCN/NH_4OAc_{aq}$ (0.1%) to 95% $MeCN//NH_4OAc_{aq}$ (0.1%)) gave a gum. The product was dissolved in methanol and treated with 1.0M ethereal Hydrogen chloride solution to give the product as a solid (0.045 g).

MS: ESI 465 (M+H)

^{1}H NMR: δ(DMSO) 8.58-8.63 (m, 1H), 7.01-7.58 (m, 6H), 4.80 (br s, 1H), 4.23-4.59 (m, 1H), 3.83 (s, 3H), 3.04-3.60 (m, 4H), 1.89-2.14 (m, 5H), 0.85 (m, 6H)

Example 29

N-{2-[4-(3,4-Dichloroanilino)-1-piperidinyl]ethyl}-3-methoxybenzamide dihydrochloride



(i) tert-Butyl 4-(3,4-dichloroanilino)-1-piperidinecarboxylate

A solution of 3,4-dichloroaniline (5g), *N*-tert-butoxycarbonyl-4-piperidone (11.7g), sodium triacetoxyborohydride (19.7g) and acetic acid (7ml) in dichloroethane (150ml) was stirred for 16 hours. 2M NaOH solution and ether were added, the organic phase separated, dried and concentrated. The residue was triturated under an isohexane : ethyl acetate, 4:1 mixture and the sub-titled product collected as a solid (7.25g).

MS: APCI(+ve) 345 (M+H)

^{1}H NMR: δ(DMSO) 7.23 (d,1H), 6.77 (d,1H), 6.57 (dd, 1H), 5.99 (d,1H), 3.85 (bd, 2H), 3.40 (m, 1H), 2.90 (bm, 2H), 1.85 (m, 2H), 1.39 (s, 9H), 1.19 (m,2H)

(ii) N-(3,4-Dichlorophenyl)-4-piperidinamine trifluoroacetate

The product of step (i) above (6.5g) was dissolved in dichloromethane (75ml) and trifluoroacetic acid (25ml) added. After 72 hours at room temperature the solution was evaporated and the residue triturated under ether to give the sub-titled product as a solid (6.3g).

MS: APCI(+ve) 245/7 (M+H)

^{1}H NMR: δ(DMSO) 8.65 (bs, 1H), 8.50 (bs, 1H), 7.26 (d, 1H), 6.81 (d, 1H), 6.60 (dd, 1H), 6.19 (bs, 1H), 3.53 (bs, 1H), 3.30 (m, 2H), 3.0 (m, 2H), 2.02 (m, 2H), 1.50 (m, 2H)

(iii) tert-Butyl 2-[4-(3,4-dichloroanilino)-1-piperidinyl]ethylcarbamate

The product from step (ii) above (2.0g), N-tert-butoxycarbonyl-2-bromoethanamine (1.0g) and N,N-di-isopropylethylamine (3.7ml) were dissolved in DMF (25ml) and stirred for 16 hours. Water and ethyl acetate were added, the organic phase separated, dried and evaporated to give a gum. Purification by chromatography (dichloromethane : methanol, 95:5) gave the sub-titled product as a solid (1.25g).

MS: APCI(+ve) 388/90 (M+H)

^{1}H NMR: δ(DMSO) 7.22 (d, 1H), 6.73 (d, 1H), 6.62 (t,1H), 6.54 (dd, 1H) 5.94 (d, 1H), 3.17 (m, 1H), 3.02 (m, 2H), 2.77 (bd, 2H), 2.31 (t, 3H), 2.06 (t, 2H), 1.84 (bd, 2H), 1.35 (m, 11H)

(iv) 1-(2-Aminoethyl)-N-(3,4-dichlorophenyl)-4-piperidinamine triluoroacetate

The product from step (iii) above (1.2g) was dissolved in dichloromethane (30ml) and trifluoroacetic acid (10ml) added. After 72 hours at room temperature the reaction mixture

was evaporated and residue triturated under ether to give the sub-titled product as a solid (1.6g).

MS: APCI(+ve) 288/90 (M+H)

(v) N-{2-[4-(3,4-Dichloroanilino)-1-piperidinyl]ethyl}-3-methoxybenzamide dihydrochloride

The product of step (iv) above (0.5g) and triethylamine (1.1ml) were dissolved in DMF (10ml), 3-methoxybenzoylchloride (0.11ml) was added dropwise. After 2 hours, water and ethyl acetate were added, the organic phase separated, dried and evaporated. Purification of the residue by chromatography (dichloromethane : methanol, 95:5) gave an oil which was treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.15g).

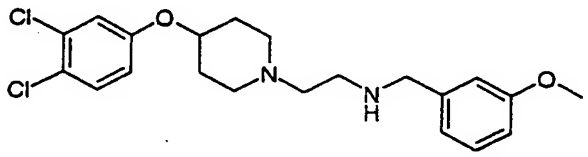
MS: ESI 422.14 (M+H)

^{1}H NMR: δ(DMSO) 10.44 (bs, 1H), 8.93 (t, 1H) 7.51 (m, 2H), 7.40 (t, 1H), 7.26 (d,1H), 7.11 (dd, 1H), 6.81 (d, 1H), 6.60 (dd, 1H), 3.82 (s, 3H), 2.68 (m, 4H), 3.25 (m, 5H), 2.09 (bd, 2H), 1.76 (m, 2H)

Melting point: 170 °C

Example 30

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N-(3-methoxybenzyl)amine dihydrochloride



A suspension of the product of Example 1 step (iv) (0.11g) in a mixture of DMF (1.5ml) and 1,2-dichloroethane (3ml) was stirred under an atmosphere of nitrogen. Sodium triacetoxyborohydride (0.097g), 3-methoxybenzaldehyde (0.041g) and triethylamine (0.046g) were added and the mixture stirred for 18 hours at room temperature. Chloroform

and aqueous $NaHCO_3$ solution were added, the organic phase separated, dried and concentrated to a gum. Purification by chromatography (chloroform : triethylamine : methanol, 89 :10:1) gave an oil which was treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.067g).

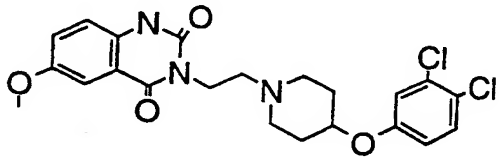
MS: ESI 409.14 (M+H)

^{1}H NMR: δ(DMSO) 7.50 (d, 1H), 7.30 (m, 3H), 7.12 (d, 1H), 7.03 (dd, 1H), 6.97 (dd,1H), 4.71 (bm, 1H), 4.18 (s, 2H), 3.80 (s, 3H), 3.45 (bm, 4H), 2.23 (m, 6H), 2.04 (m, 2H),

Melting point: 247-51 °C

Example 31

3-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methoxy-2,4(1H,3H)-quinazolinedione



(i) 2-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxybenzamide

Prepared by the method of Example 2 using the product from Example 1 step (iv) (1.0g) and 2-amino-5-methoxybenzoic acid (0.418g) without the addition of 1.0M ethereal hydrogen chloride solution to give an oil which was purified by chromatography (dichloromethane : methanol, 95:5) to give the sub-titled product as an oil (0.82g).

MS: APCI(+ve) 438 (M+H)

(ii) 3-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methoxy-2,4(1H,3H)-quinazolinedione

The product of step (i) above was dissolved in toluene (10ml). A solution of phosgene 2.0M in toluene (10ml) was added, the solution heated under reflux for 1 hour and cooled. Ethyl acetate and aqueous $NaHCO_3$ solution were added, the organic phase separated,

dried and concentrated to leave a residue which was purified by chromatography (dichloromethane : methanol, 95:5). The titled product was obtained as a solid (0.11g).

MS: ESI 464.11 (M+H)

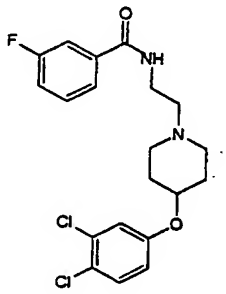
^{1}H NMR: δ(DMSO) 7.49 (dd, 1H), 7.36 (d, 1H), 7.30 (dd, 1H), 7.24 (d,1H), 6.98 (dd, 1H), 4.44 (m, 1H), 4.03 (t, 3H), 3.80 (s, 3H), 2.76 (m, 2H), 2.32 (m, 2H), 1.89 (m, 2H), 1.57 (m, 2H)

Melting point: 190 °C

The compounds of following Examples 32 to 125 were prepared by methods analogous to the method of Example 10.

Example 32

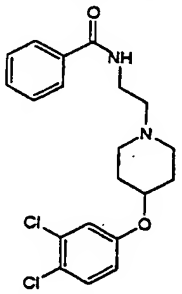
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide



MS: APC1 (+ve) BP 411

Example 33

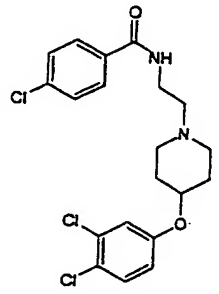
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 393

Example 34

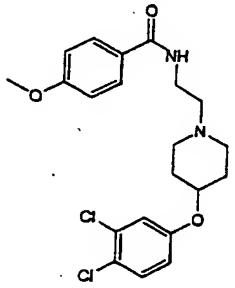
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 429

Example 35

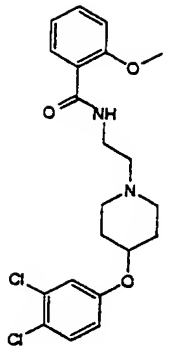
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide



MS: APC1 (+ve) BP 423

Example 36

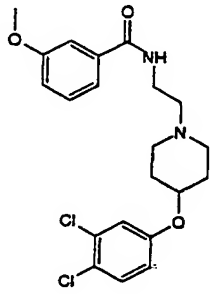
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxybenzamide



MS: APC1 (+ve) BP 423

Example 37

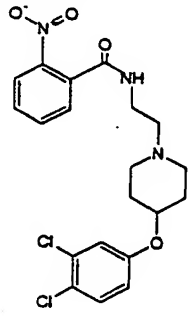
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide



MS: APC1 (+ve) BP 423

Example 38

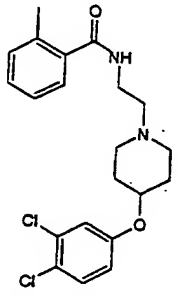
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-nitrobenzamide



MS: APC1 (+ve) BP 438

Example 39

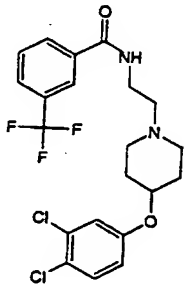
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methylbenzamide



MS: APC1 (+ve) BP 407

Example 40

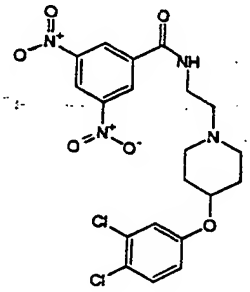
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide



MS: APC1 (+ve) BP 461

Example 41

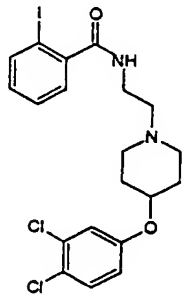
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-dinitrobenzamide



MS: APC1 (+ve) BP 483

Example 42

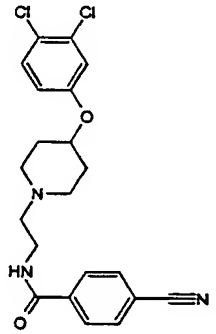
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-iodobenzamide



MS: APC1 (+ve) BP 519

Example 43

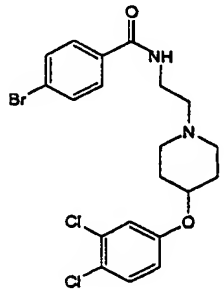
4-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 418

Example 44

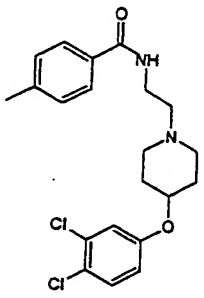
4-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 473

Example 45

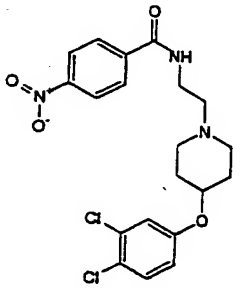
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methylbenzamide



MS: APC1 (+ve) BP 407

Example 46

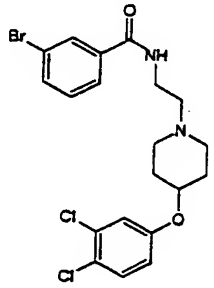
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-nitrobenzamide



MS: APC1 (+ve) BP 438

Example 47

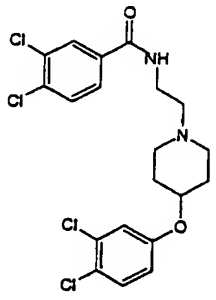
3-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 473

Example 48

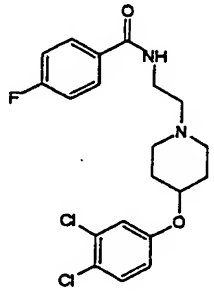
3,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 463

Example 49

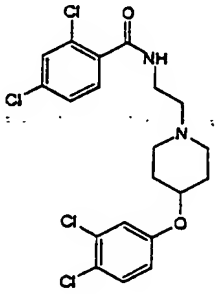
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide



MS: APC1 (+ve) BP 411

Example 50

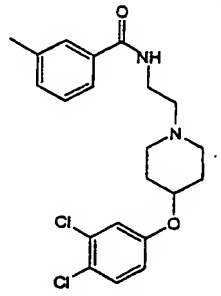
2,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 463

Example 51

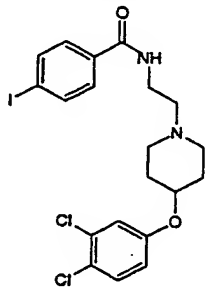
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methylbenzamide



MS: APC1 (+ve) BP 407

Example 52

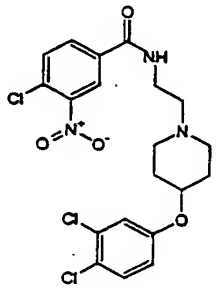
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-iodobenzamide



MS: APC1 (+ve) BP 519

Example 53

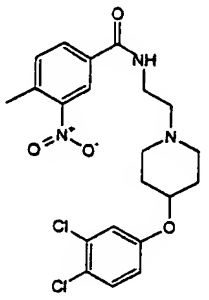
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-nitrobenzamide



MS: APC1 (+ve) BP 472

Example 54

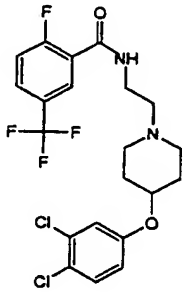
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide



MS: APC1 (+ve) BP 452

Example 55

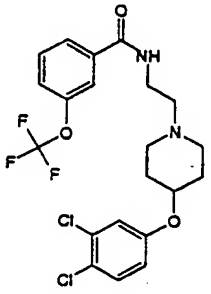
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-5-(trifluoromethyl)benzamide



MS: APC1 (+ve) BP 479

Example 56

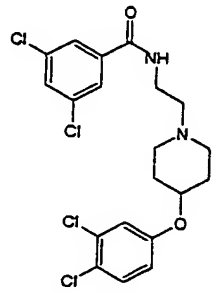
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide



MS: APC1 (+ve) BP 477

Example 57

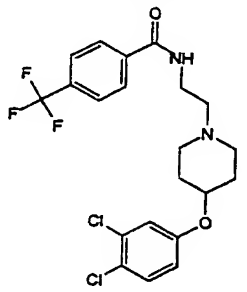
3,5-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 463

Example 58

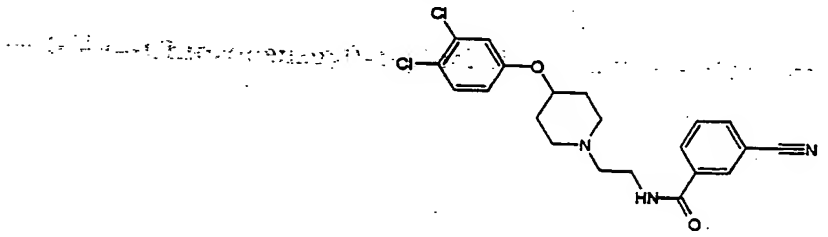
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)benzamide



MS: APC1 (+ve) BP 461

Example 59

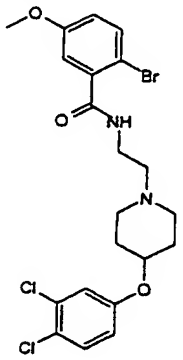
3-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 418

Example 60

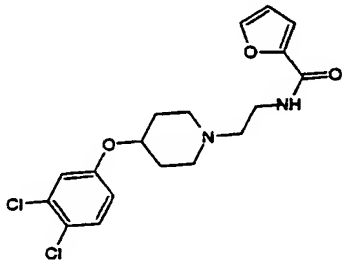
2-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxybenzamide



MS: APC1 (+ve) BP 503

Example 61

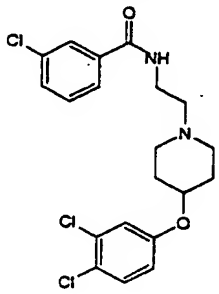
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide



MS: APC1 (+ve) BP 383

Example 62

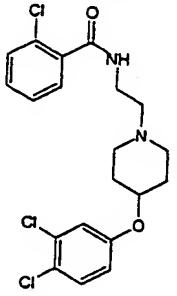
3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 427

Example 63

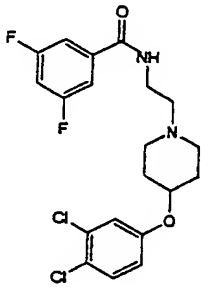
2-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 429

Example 64

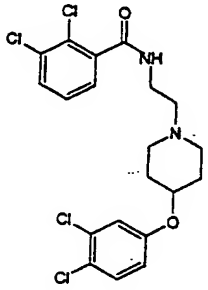
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-difluorobenzamide



MS: APC1 (+ve) BP 429

Example 65

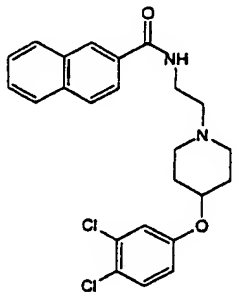
2,3-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 463

Example 66

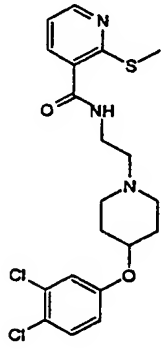
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-naphthamide



MS: APC1 (+ve) BP 442

Example 67

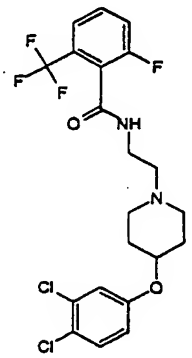
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide



MS: APC1 (+ve) BP 440

Example 68

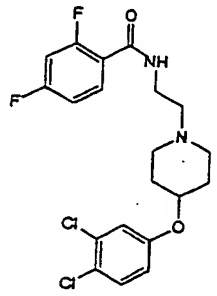
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-6-(trifluoromethyl)benzamide



MS: APC1 (+ve) BP 479

Example 69

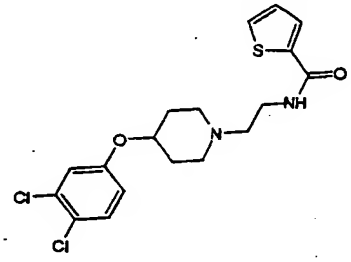
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-difluorobenzamide



MS: APC1 (+ve) BP 429

Example 70

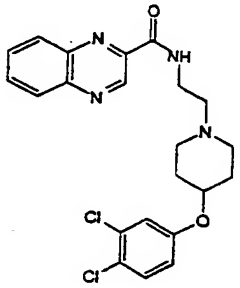
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-thiophenecarboxamide



MS: APC1 (+ve) BP 399

Example 71

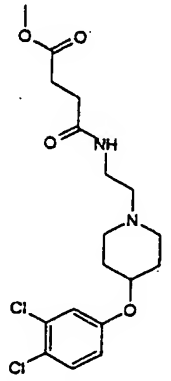
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide



MS: APC1 (+ve) BP 445

Example 72

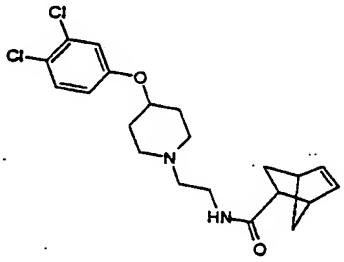
Methyl 4-({2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}amino)-4-oxobutanoate



MS: APC1 (+ve) BP 403

Example 73

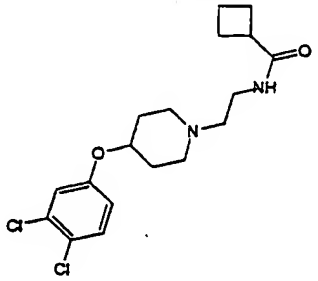
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide



MS: APC1 (+ve) BP 409

Example 74

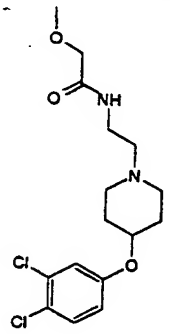
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclobutanecarboxamide



MS: APC1 (+ve) BP 371

Example 75

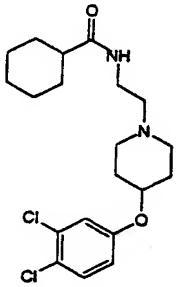
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxyacetamide



MS: APC1 (+ve) BP 361

Example 76

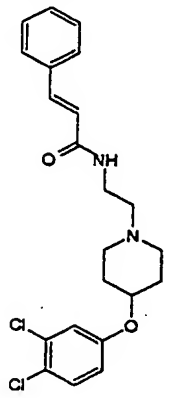
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclohexanecarboxamide



MS: APC1 (+ve) BP 399

Example 77

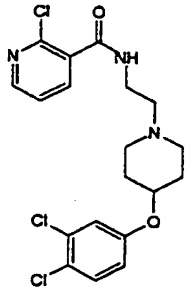
(E)-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide



MS: APC1 (+ve) BP 419

Example 78

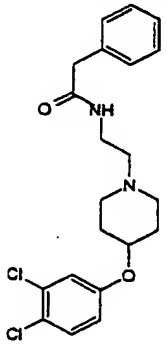
2-Chloro-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}nicotinamide



MS: APC1 (+ve) BP 430

Example 79

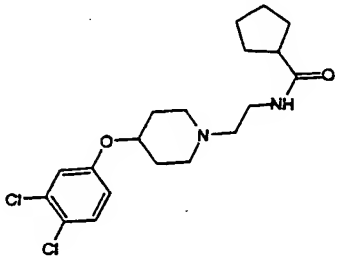
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide



MS: APC1 (+ve) BP 407

Example 80

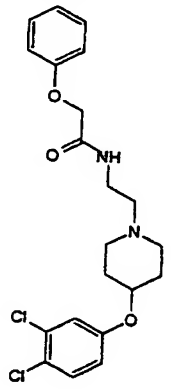
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclopentanecarboxamide



MS: APC1 (+ve) BP 385

Example 81

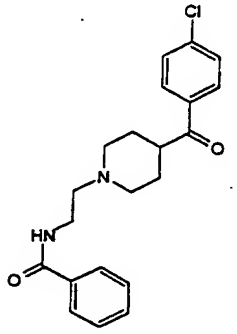
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenoxyacetamide



MS: APC1 (+ve) BP 423

Example 82

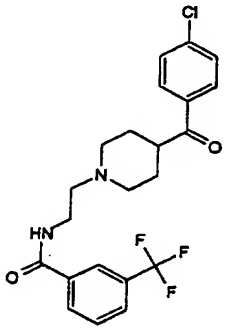
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}benzamide



MS: APC1 (+ve) BP 371

Example 83

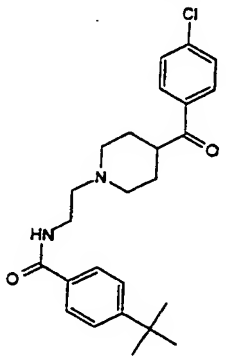
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide



MS: APC1 (+ve) BP 439

Example 84

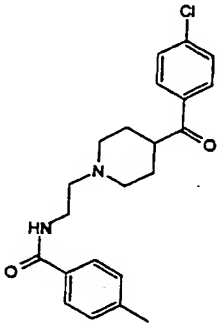
4-(tert-Butyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}benzamide



MS: APCI (+ve) BP 427

Example 85

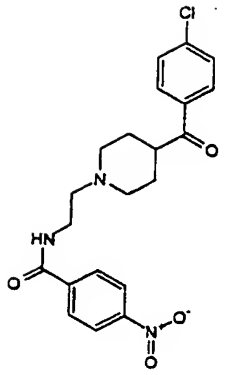
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methylbenzamide



MS: APCI (+ve) BP 385

Example 86

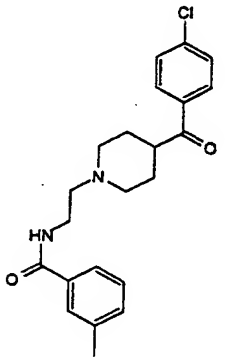
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-nitrobenzamide



MS: APC1 (+ve) BP 416

Example 87

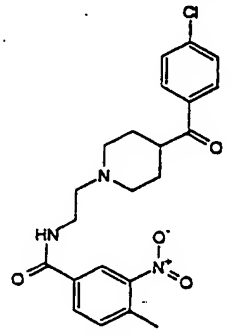
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-methylbenzamide



MS: APC1 (+ve) BP 385

Example 88

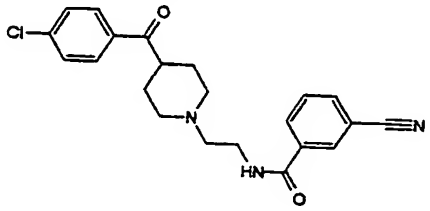
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide



MS: APC1 (+ve) BP 430

Example 89

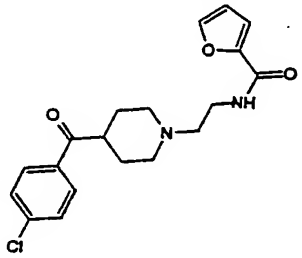
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-cyanobenzamide



MS: APC1 (+ve) BP 396

Example 90

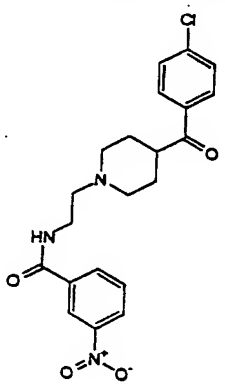
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-furamide



MS: APC1 (+ve) BP 361

Example 91

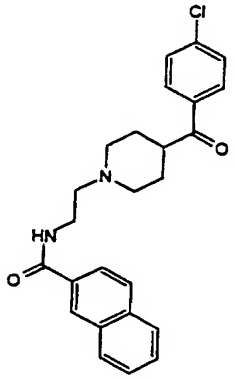
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-nitrobenzamide



MS: APC1 (+ve) BP 416

Example 92

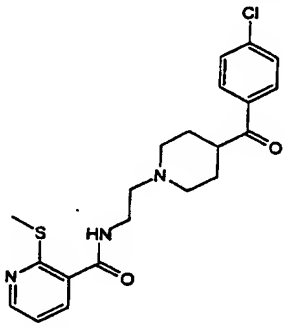
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-naphthamide



MS: APC1 (+ve) BP 421

Example 93

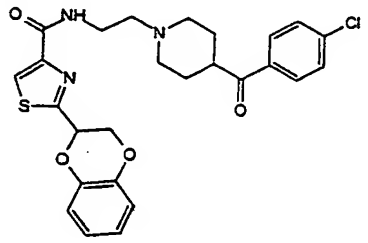
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide



MS: APC1 (+ve) BP 418

Example 94

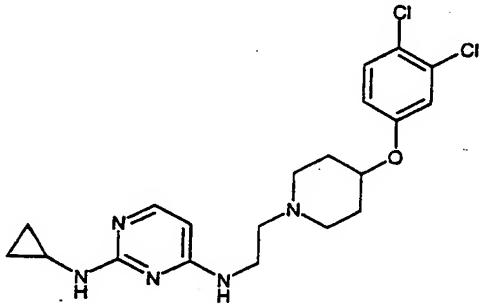
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(2,3-dihydro-1,4-benzodioxin-2-yl)-1,3-thiazole-4-carboxamide



MS: APC1 (+ve) BP 512

Example 95

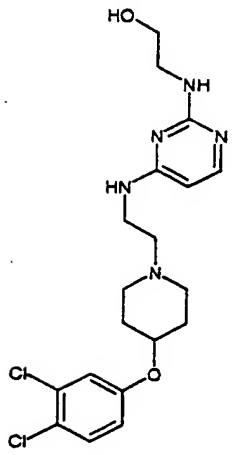
N~2~-Cyclopropyl-N~4~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine



MS: APCI(+ve) 422 (M+1)

Example 96

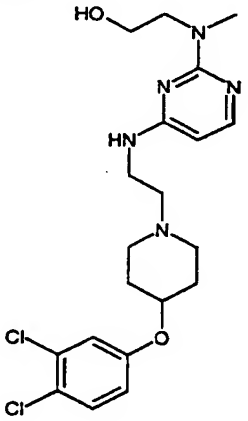
2-{[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl]amino}-1-ethanol



MS: APCI(+ve) 426 (M+1)

Example 97

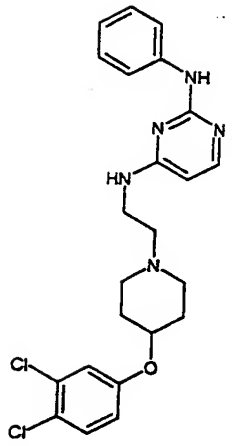
2-[[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol



MS: APCI(+ve) 440 (M+1)

Example 98

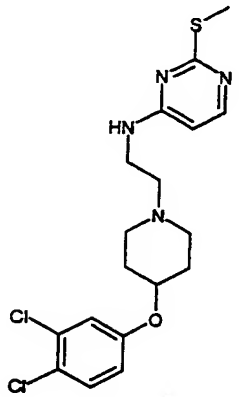
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~-phenyl-2,4-pyrimidinediamine



MS: APCI(+ve) 458 (M+1)

Example 99

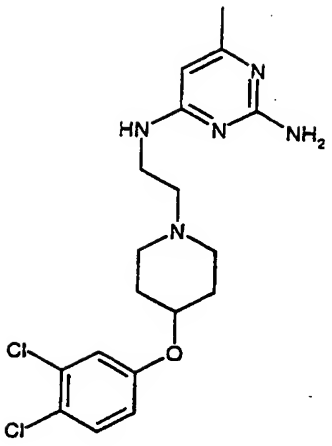
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 413 (M+1)

Example 100

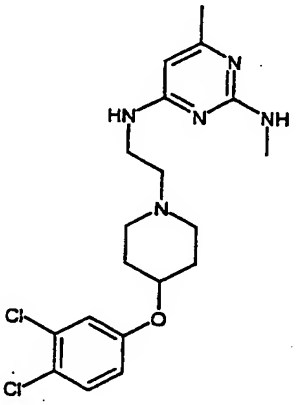
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2,4-pyrimidinediamine



MS: APCI(+ve) 396 (M+1)

Example 101

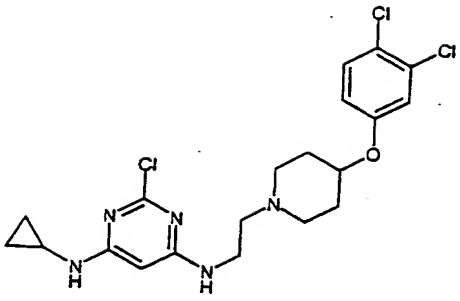
N~4~~{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 410 (M+1)

Example 102

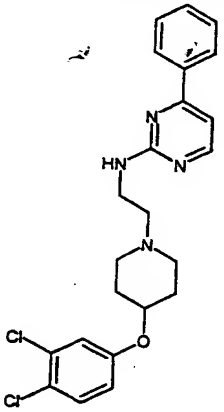
2-Chloro-N~4~~cyclopropyl-N~6~~{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-pyrimidinediamine



MS: APCI(+ve) 456 (M+1)

Example 103

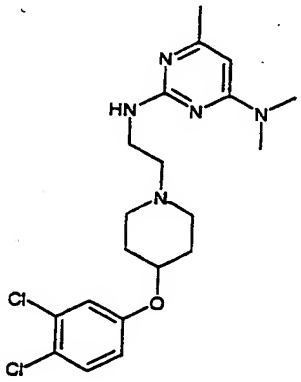
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenyl-2-pyrimidinamine



MS: APCI(+ve) 443 (M+1)

Example 104

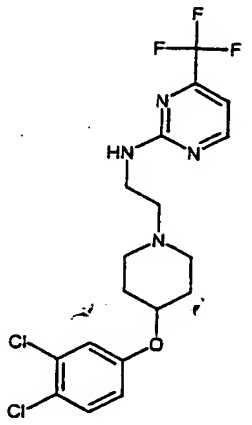
N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 424 (M+1)

Example 105

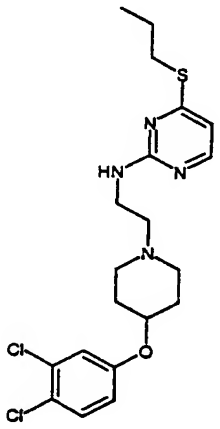
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)-2-pyrimidinamine



MS: APCI(+ve) 435 (M+1)

Example 106

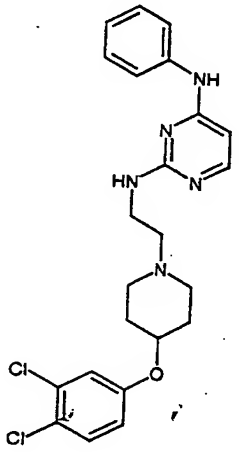
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(propylsulfanyl)-2-pyrimidinamine



MS: APCI(+ve) 441 (M+1)

Example 107

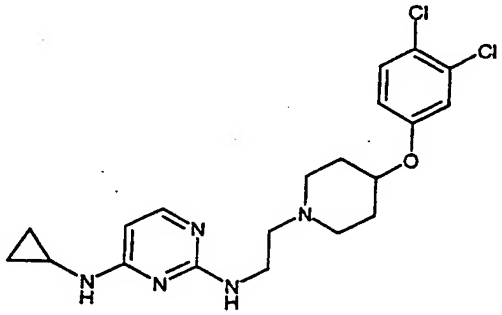
N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~-phenyl-2,4-pyrimidinediamine



MS: APCI(+ve) 458 (M+1)

Example 108

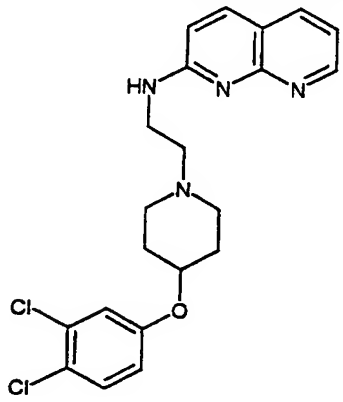
N~4~-Cyclopropyl-N~2~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine



MS: APCI(+ve) 422 (M+1)

Example 109

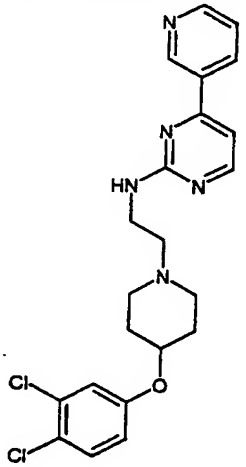
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}[1,8]naphthyridin-2-amine



MS: APCI(+ve) 417 (M+1)

Example 110

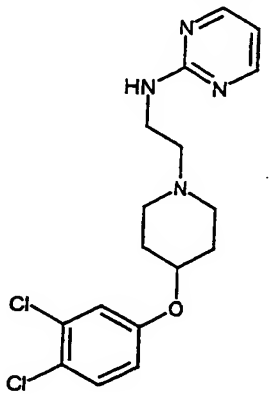
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-pyridinyl)-2-pyrimidinamine



MS: APCI(+ve) 444 (M+1)

Example 111

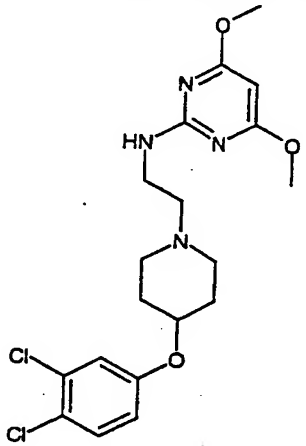
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrimidinamine



MS: APCI(+ve) 367 (M+1)

Example 112

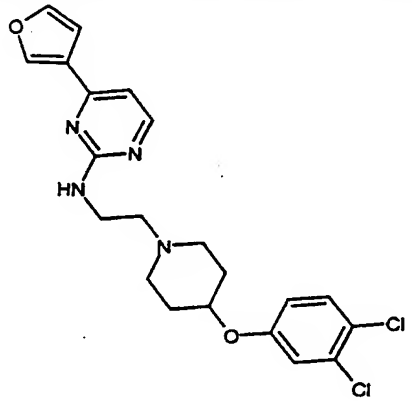
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-dimethoxy-2-pyrimidinamine



MS: APCI(+ve) 427 (M+1)

Example 113

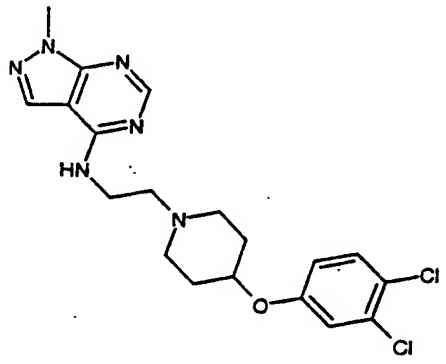
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-furyl)-2-pyrimidinamine



MS: APCI(+ve) 433 (M+1)

Example 114

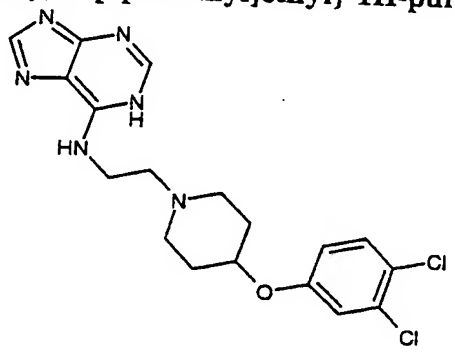
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine



MS: APCI(+ve) 421 (M+1)

Example 115

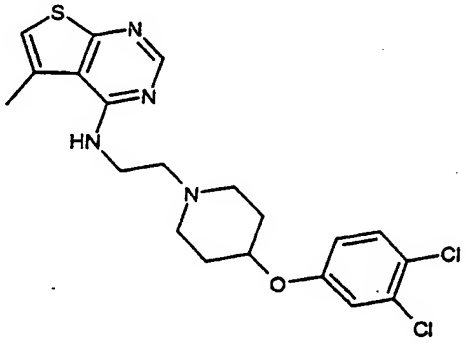
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-purin-6-amine



MS: APCI(+ve) 407 (M+1)

Example 116

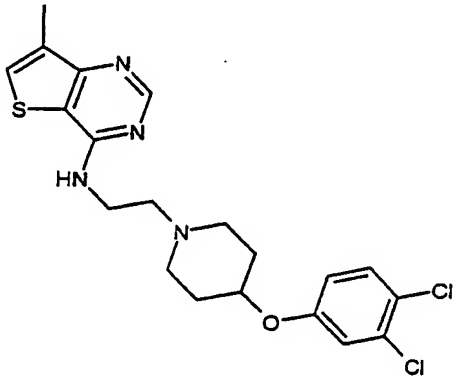
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methylthieno[2,3-d]pyrimidin-4-amine



MS: APCI(+ve) 437 (M+1)

Example 117

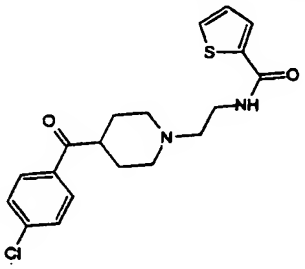
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methylthieno[3,2-d]pyrimidin-4-amine



MS: APCI(+ve) 437 (M+1)

Example 118

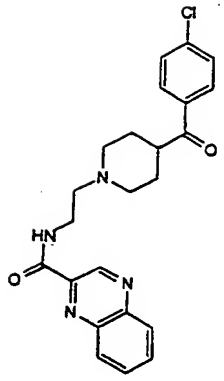
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-thiophenecarboxamide



MS: APC1 (+ve) BP 377

Example 119

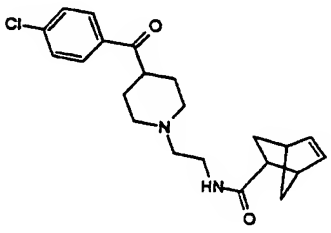
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide



MS: APC1 (+ve) BP 423

Example 120

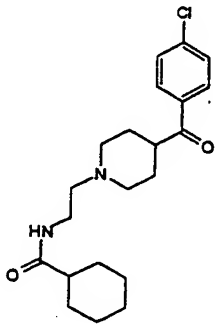
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide



MS: APC1 (+ve) BP 387

Example 121

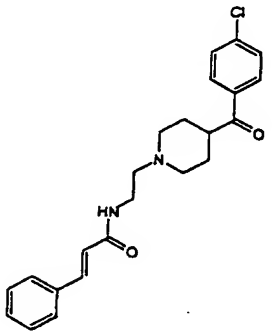
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}cyclohexanecarboxamide



MS: APC1 (+ve) BP 377

Example 122

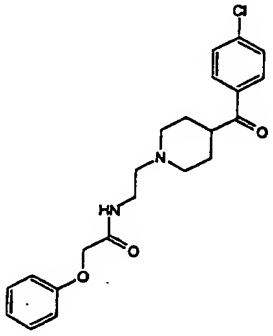
(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide



MS: APC1 (+ve) BP 397

Example 123

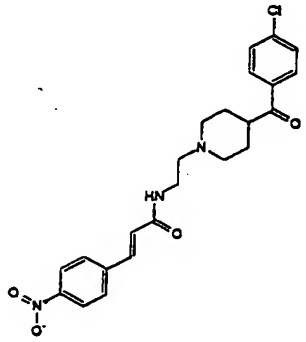
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-phenoxyacetamide



MS: APC1 (+ve) BP 401

Example 124

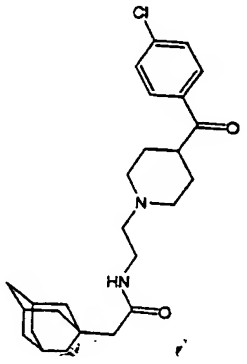
(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(4-nitrophenyl)-2-propenamide



MS: APCI (+ve) BP 442

Example 125

2-(1-Adamantyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}acetamide

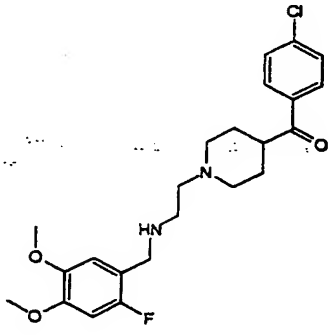


MS: APCI (+ve) BP 443

The compounds of following Examples 126 to 168 were prepared by methods analogous to the method of Example 30.

Example 126

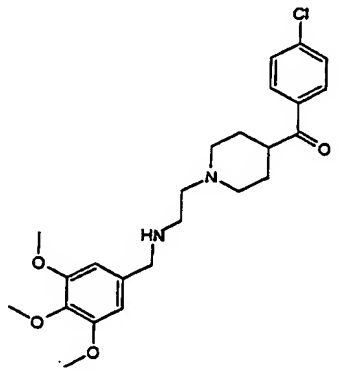
(4-Chlorophenyl)(1-{2-[(2-fluoro-4,5-dimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APCI (+ve) BP 435

Example 127

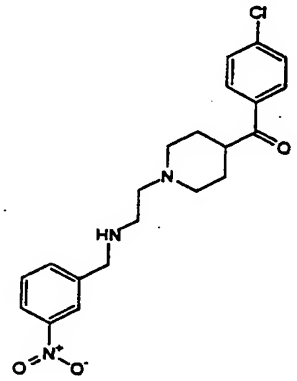
(4-Chlorophenyl)(1-{2-[(3,4,5-trimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 447

Example 128

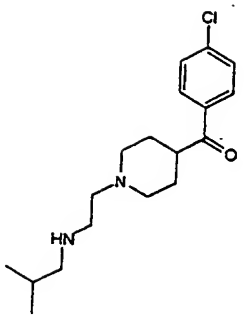
(4-Chlorophenyl)(1-{2-[(3-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 402

Example 129

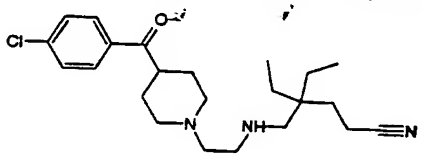
(4-Chlorophenyl){1-[2-(isobutylamino)ethyl]-4-piperidinyl}methanone



MS: APCI (+ve) BP 323

Example 130

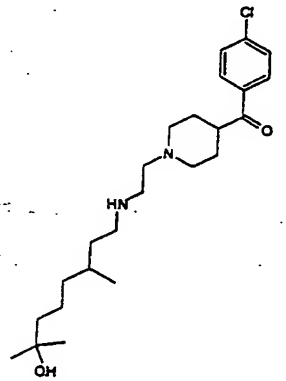
4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-ethylhexanenitrile



MS: APCI (+ve) BP 404

Example 131

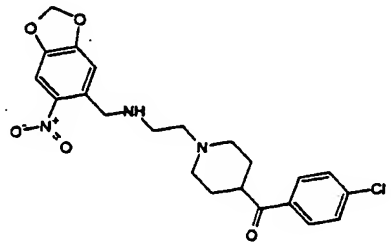
(4-Chlorophenyl)(1-{2-[(7-hydroxy-3,7-dimethyloctyl)amino]ethyl}-4-piperidinyl)methanone



MS: APCI (+ve) BP 423

Example 132

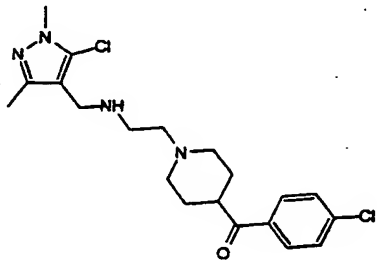
(4-Chlorophenyl)[1-(2-{[(6-nitro-1,3-benzodioxol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 446

Example 133

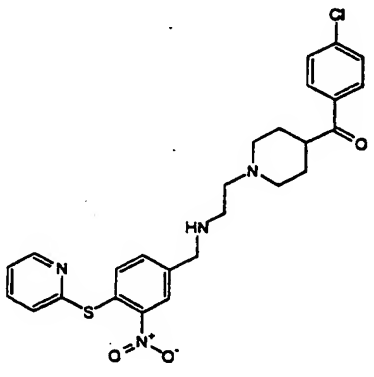
[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone



MS: APC1 (+ve) BP 409

Example 134

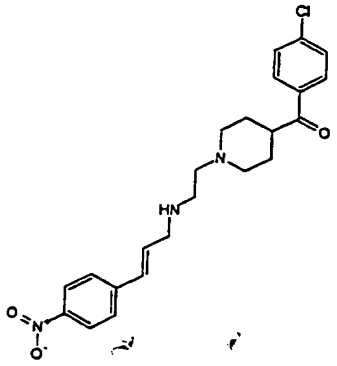
(4-Chlorophenyl)[1-(2-{[3-nitro-4-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 511

Example 135

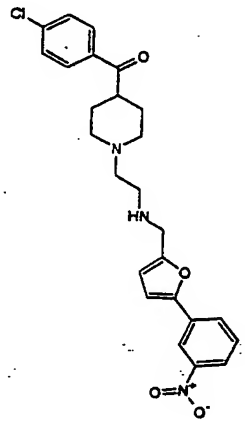
(4-Chlorophenyl)[1-(2-{[(E)-3-(4-nitrophenyl)-2-propenyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 428

Example 136

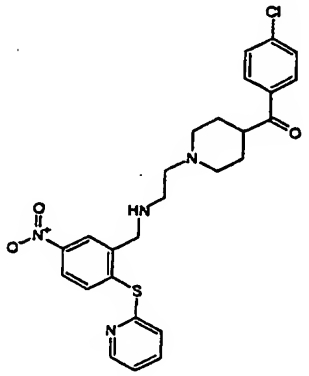
(4-Chlorophenyl){1-[2-({[5-(3-nitrophenyl)-2-furyl]methyl}amino)ethyl]-4-piperidinyl}methanone



MS: APC1 (+ve) BP 468

Example 137

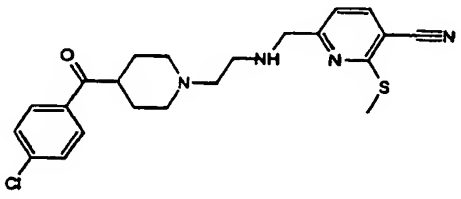
(4-Chlorophenyl)[1-(2-{[5-nitro-2-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 511

Example 138

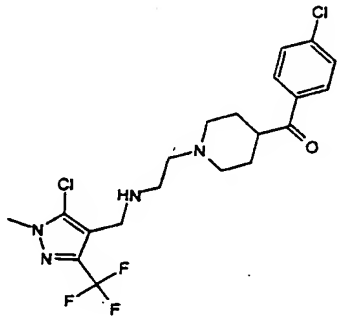
6-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-2-(methylsulfanyl)nicotinonitrile



MS: APC1 (+ve) BP 429

Example 139

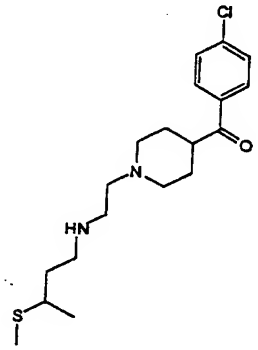
{1-[2-({[5-Chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazol-4-yl]methyl}amino)ethyl]-4-piperidinyl}(4-chlorophenyl)methanone



MS: APC1 (+ve) BP 463

Example 140

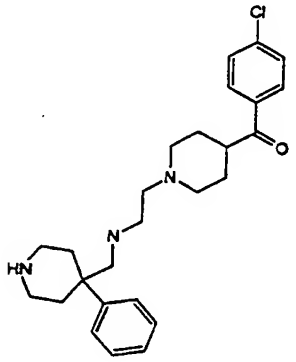
(4-Chlorophenyl)[1-(2-{[3-(methylsulfanyl)butyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 369

Example 141

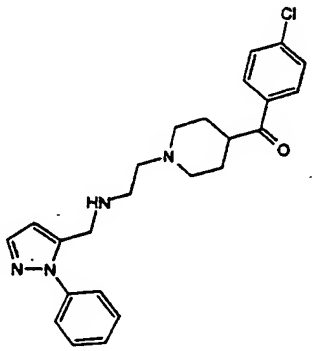
(4-Chlorophenyl)[1-(2-{[(4-phenyl-4-piperidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 440

Example 142

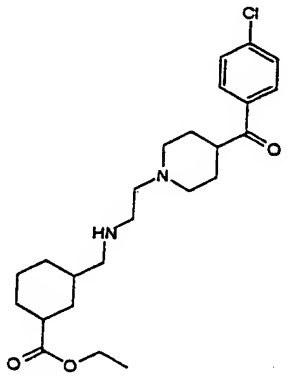
(4-Chlorophenyl)[1-(2-{[(1-phenyl-1H-pyrazol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 423

Example 143

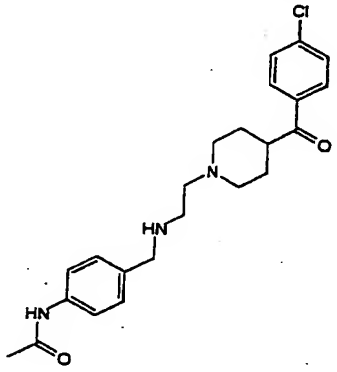
Ethyl 3-[({2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}-amino)methyl]cyclohexanecarboxylate



MS: APCI (+ve) BP 435

Example 144

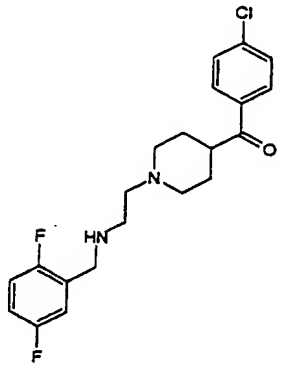
N-{4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]phenyl}acetamide



MS: APCI (+ve) BP 414

Example 145

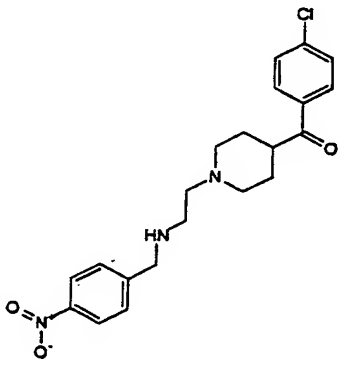
(4-Chlorophenyl)(1-{2-[(2,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 393

Example 146

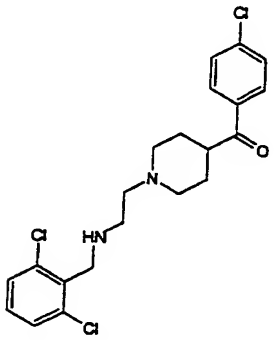
(4-Chlorophenyl)(1-{2-[(4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 402

Example 147

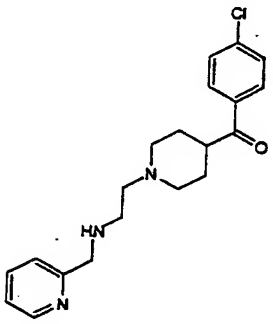
(4-Chlorophenyl)(1-{2-[(2,6-dichlorobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APCI (+ve) BP 425

Example 148

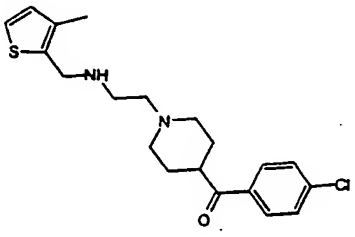
(4-Chlorophenyl)(1-{2-[(2-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone



MS: APCI (+ve) BP 358

Example 149

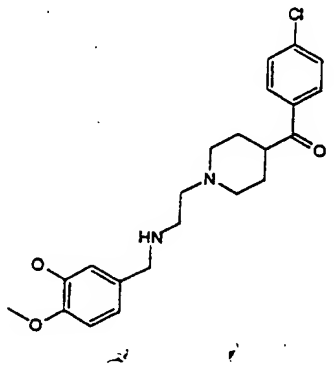
(4-Chlorophenyl)[1-(2-{[(3-methyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APCI (+ve) BP 377

Example 150

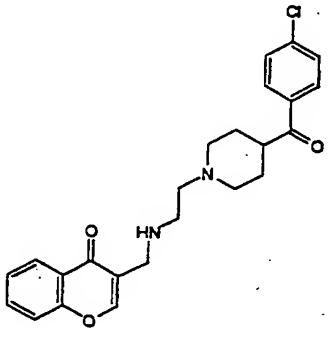
(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-methoxybenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 403

Example 151

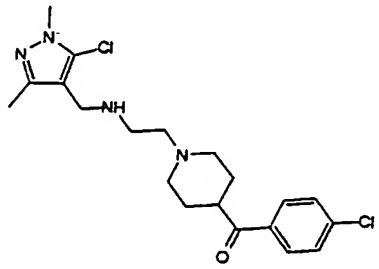
3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4H-chromen-4-one



MS: APC1 (+ve) BP 425

Example 152

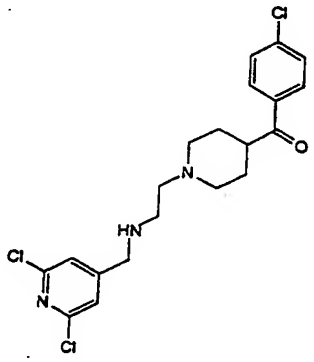
[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone



MS: APC1 (+ve) BP 409

Example 153

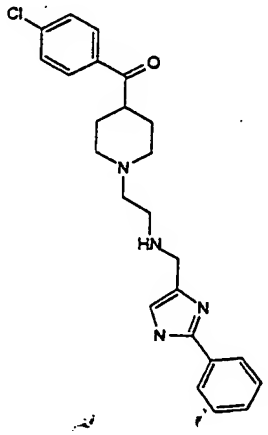
(4-Chlorophenyl)[1-(2-{[(2,6-dichloro-4-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 428

Example 154

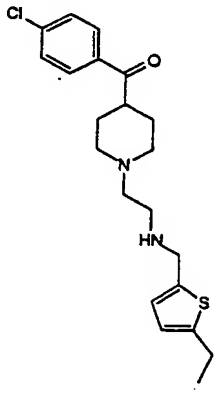
(4-Chlorophenyl)[1-(2-{[(2-phenyl-1H-imidazol-4-yl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 423

Example 155

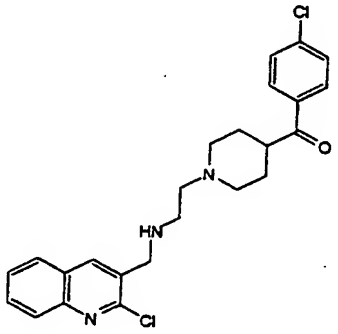
(4-Chlorophenyl)[1-(2-{[(5-ethyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 391

Example 156

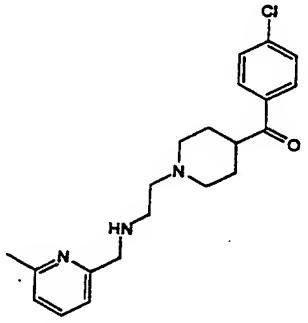
(4-Chlorophenyl)[1-(2-{[(2-chloro-3-quinolinyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APCI (+ve) BP 442

Example 157

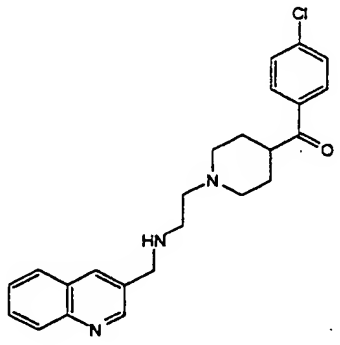
(4-Chlorophenyl)[1-(2-{[(6-methyl-2-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APCI (+ve) BP 372

Example 158

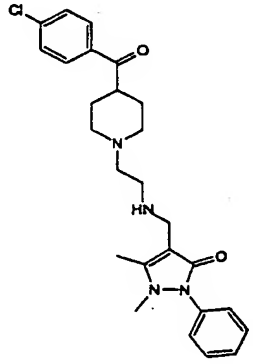
(4-Chlorophenyl)(1-{2-[(3-quinolinylmethyl)amino]ethyl}-4-piperidinyl)methanone



MS: APCI (+ve) BP 408

Example 159

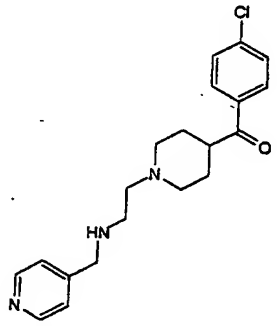
4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-1,5-dimethyl-2-phenyl-1,2-dihydro-3H-pyrazol-3-one



MS: APCI (+ve) BP 467

Example 160

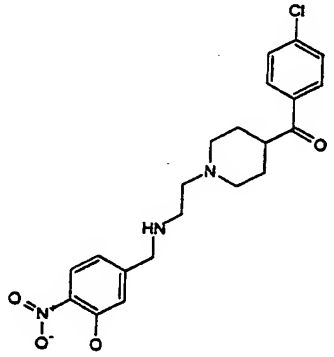
(4-Chlorophenyl)(1-{2-[(4-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 358

Example 161

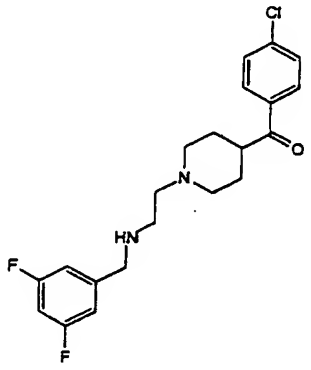
(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 418

Example 162

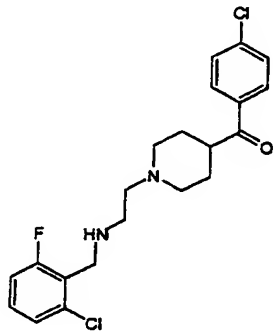
(4-Chlorophenyl)(1-{2-[(3,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone



MS: APC1 (+ve) BP 393

Example 163

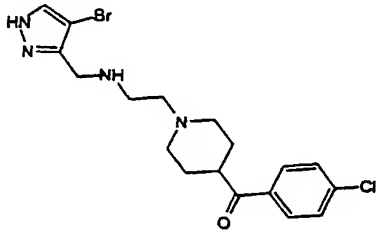
(1-{2-[(2-Chloro-6-fluorobenzyl)amino]ethyl}-4-piperidinyl)(4-chlorophenyl)methanone



MS: APC1 (+ve) BP 409

Example 164

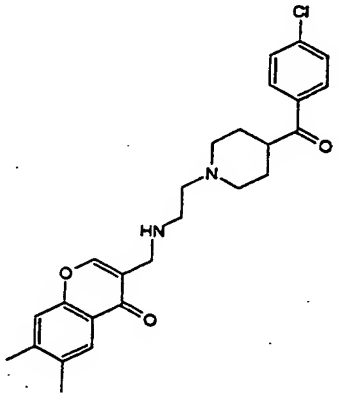
[1-(2-{[(4-Bromo-1H-pyrazol-3-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone



MS: APCI (+ve) BP 427

Example 165

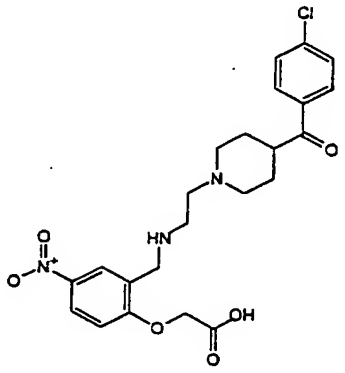
3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-6,7-dimethyl-4H-chromen-4-one



MS: APCI (+ve) BP 453

Example 166

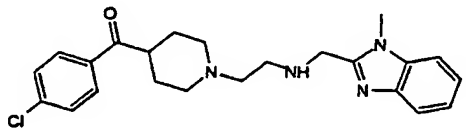
2-{2-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-nitrophenoxy}acetic acid



MS: APC1 (+ve) BP 476

Example 167

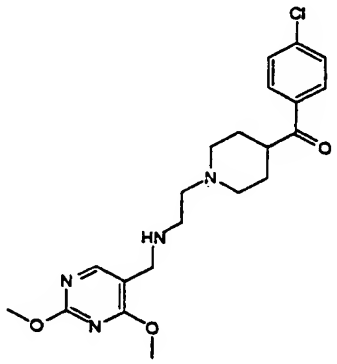
(4-Chlorophenyl)[1-(2-{[(1-methyl-1H-benzimidazol-2-yl)methyl]amino}ethyl)-4-piperidinyl]methanone



MS: APC1 (+ve) BP 411

Example 168

(4-Chlorophenyl)[1-(2-{[(2,4-dimethoxy-5-pyrimidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone

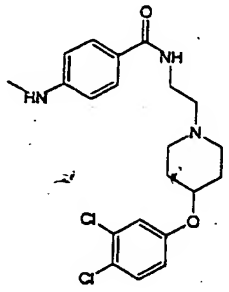


MS: APC1 (+ve) BP 419

The compounds of following Examples 169 to 209 were prepared by methods analogous to the method of Example 2.

Example 169

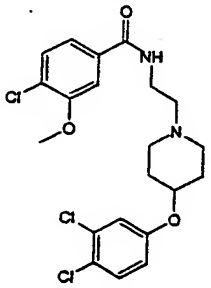
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(methylamino)benzamide



MS: APC1 (+ve) BP 422

Example 170

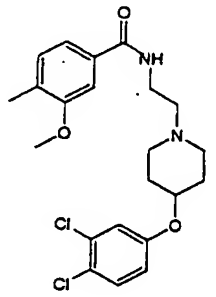
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide



MS: APC1 (+ve) BP 459

Example 171

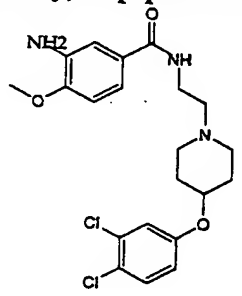
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxy-4-methylbenzamide



MS: APC1 (+ve) BP 437

Example 172

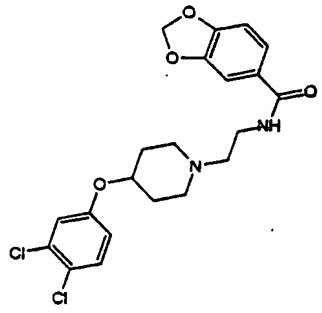
3-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide



MS: APC1 (+ve) BP 438

Example 173

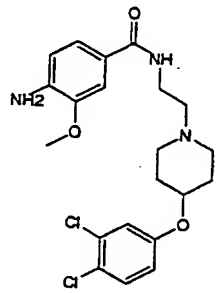
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzodioxole-5-carboxamide



MS: APC1 (+ve) BP 437

Example 174

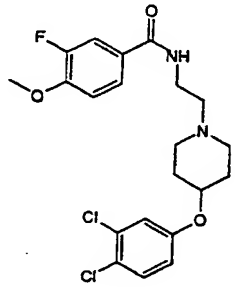
4-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide



MS: APC1 (+ve) BP 438

Example 175

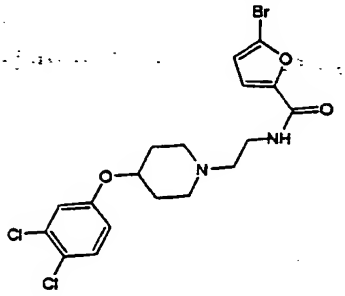
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluoro-4-methoxybenzamide



MS: APC1 (+ve) BP 441

Example 176

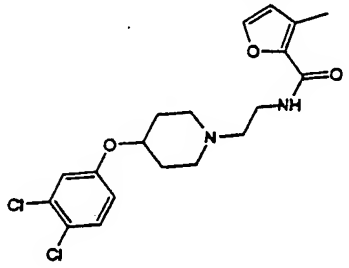
5-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide



MS: APC1 (+ve) BP 463

Example 177

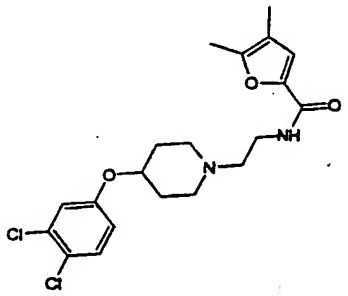
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methyl-2-furamide



MS: APC1 (+ve) BP 397

Example 178

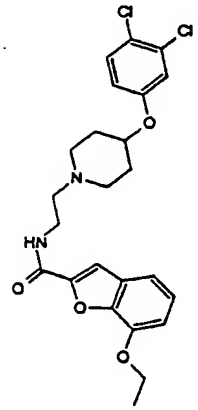
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,5-dimethyl-2-furamide



MS: APC1 (+ve) BP 411

Example 179

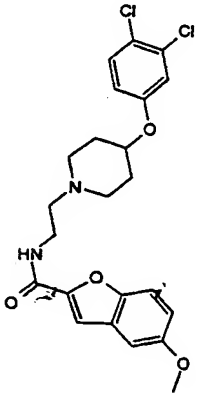
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-ethoxy-1-benzofuran-2-carboxamide



MS: APC1 (+ve) BP 477

Example 180

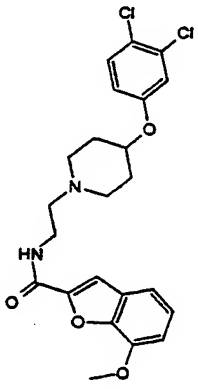
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-1-benzofuran-2-carboxamide



MS: APC1 (+ve) BP 463

Example 181

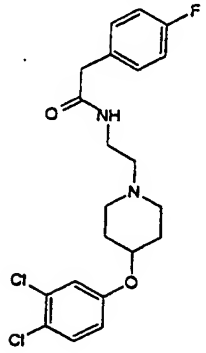
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methoxy-1-benzofuran-2-carboxamide



MS: APC1 (+ve) BP463

Example 182

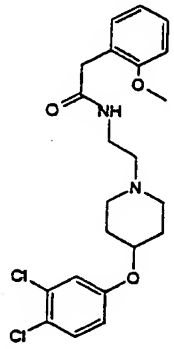
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-fluorophenyl)acetamide



MS: APC1 (+ve) BP 425

Example 183

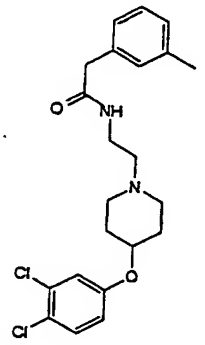
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methoxyphenyl)acetamide



MS: APC1 (+ve) BP 437

Example 184

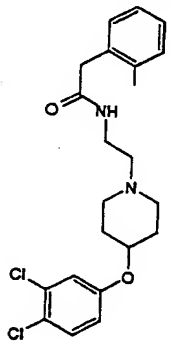
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methylphenyl)acetamide



MS: APC1 (+ve) BP421

Example 185

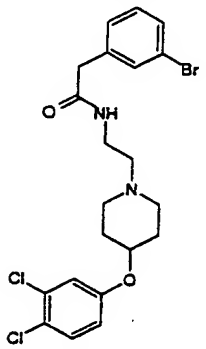
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methylphenyl)acetamide



MS: APC1 (+ve) BP 421

Example 186

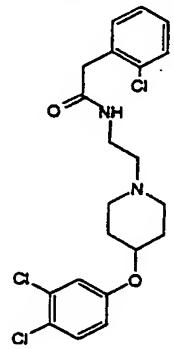
2-(3-Bromophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide



MS: APC1 (+ve) BP 487

Example 187

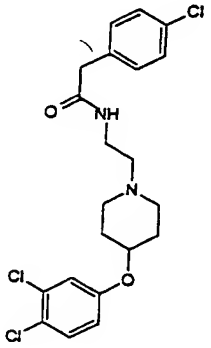
2-(2-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide



MS: APC1 (+ve) BP 441

Example 188

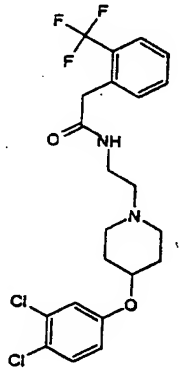
2-(4-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide



MS: APCI (+ve) BP 443

Example 189

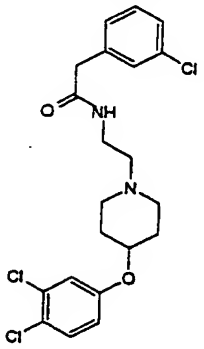
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(trifluoromethyl)phenyl]acetamide



MS: APCI (+ve) BP 475

Example 190

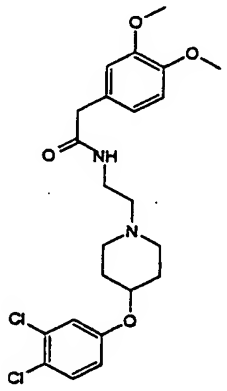
2-(3-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide



MS: APC1 (+ve) BP441

Example 191

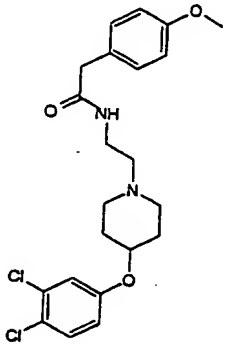
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dimethoxyphenyl)acetamide



MS: APC1 (+ve) BP467

Example 192

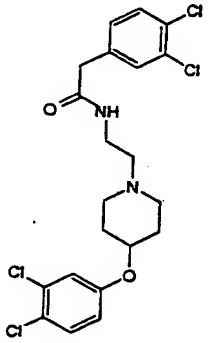
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methoxyphenyl)acetamide



MS: APC1 (+ve) BP 437

Example 193

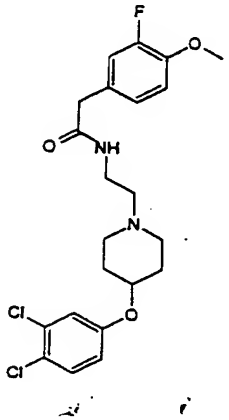
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dichlorophenyl)acetamide



MS: APC1 (+ve) BP 477

Example 194

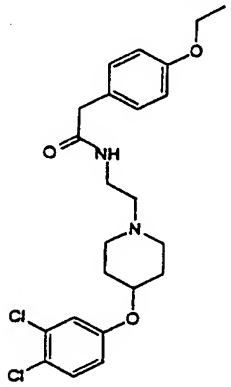
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-fluoro-4-methoxyphenyl)acetamide



MS: APC1 (+ve) BP 455

Example 195

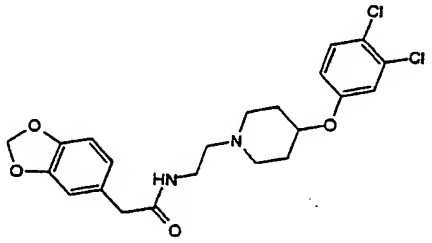
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-ethoxyphenyl)acetamide



MS: APC1 (+ve) BP 451

Example 196

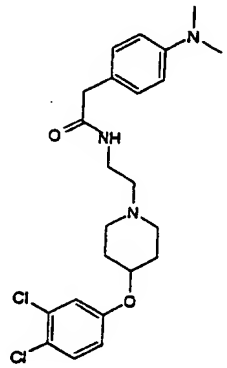
2-(1,3-Benzodioxol-5-yl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide



MS: APC1 (+ve) BP451

Example 197

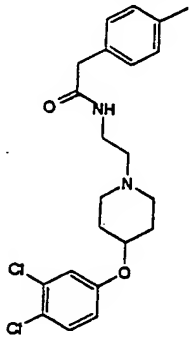
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[4-(dimethylamino)phenyl]acetamide



MS: APC1 (+ve) BP 450

Example 198

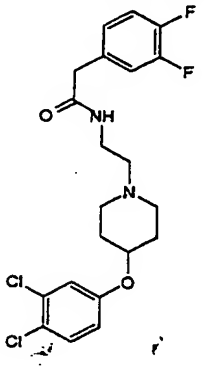
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methylphenyl)acetamide



MS: APC1 (+ve) BP 421

Example 199

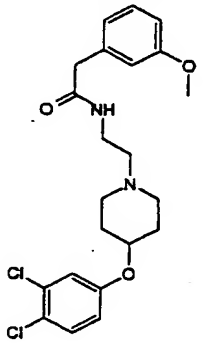
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-difluorophenyl)acetamide



MS: APC1 (+ve) BP 443

Example 200

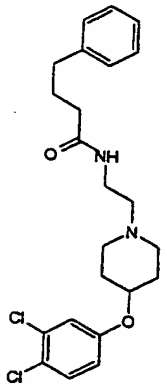
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methoxyphenyl)acetamide



MS: APC1 (+ve) BP 437

Example 201

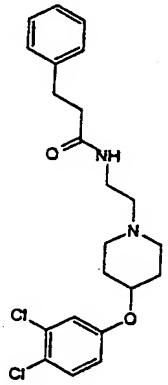
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenylbutanamide



MS: APC1 (+ve) BP 435

Example 202

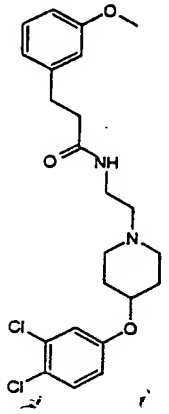
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenylpropanamide



MS: APC1 (+ve) BP 421

Example 203

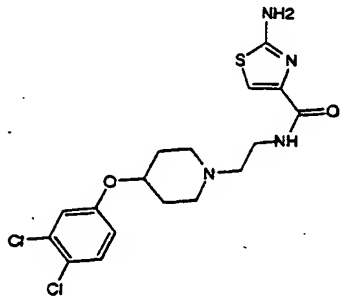
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(3-methoxyphenyl)propanamide



MS: APC1 (+ve) BP 451

Example 204

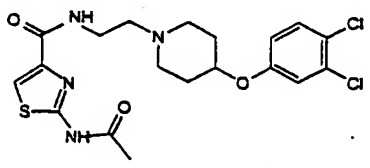
2-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide



MS: APC1 (+ve) BP 416

Example 205

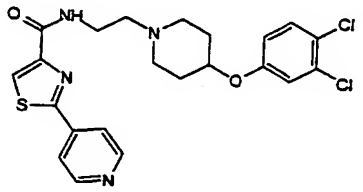
2-(Acetylamino)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide



MS: APC1 (+ve) BP 457

Example 206

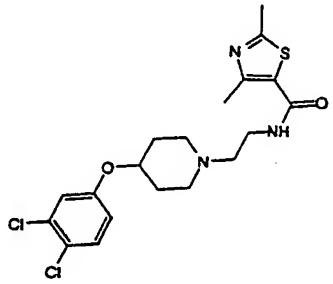
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-pyridinyl)-1,3-thiazole-4-carboxamide



MS: APC1 (+ve) BP 477

Example 207

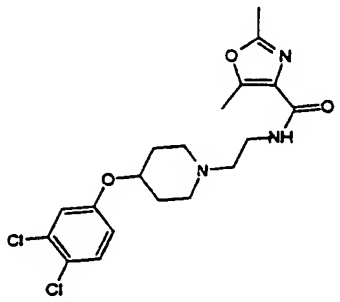
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-dimethyl-1,3-thiazole-5-carboxamide



MS: APC1 (+ve) BP 428

Example 208

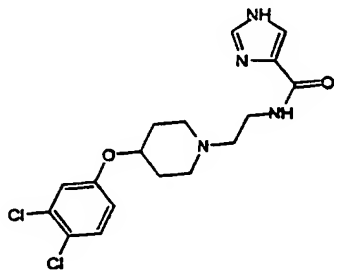
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,5-dimethyl-1,3-oxazole-4-carboxamide



MS: APCI (+ve) BP 412

Example 209

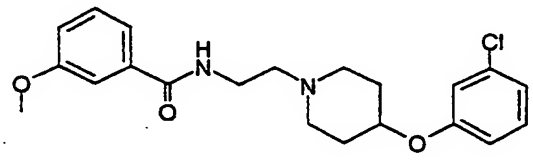
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-imidazole-4-carboxamide



MS: APCI (+ve) BP 385.

Example 210

N-{2-[4-(3,4-Chlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride



(i) 2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethylamine trifluoroacetate

Prepared by the method of Example 1 steps (i) to (iv) using 3-chlorophenol to give the product as an oil (0.5 g) which was used directly in the next step without further purification.

(ii) N-{2-[4-(3,4-Chlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride

The product of step (i) above (0.3g) was dissolved in dichloromethane (490ml), triethylamine (4 equiv) and 3-methoxybenzoyl chloride (1 equiv) were added. After 72 hours at room temperature, water was added, the organic phase separated, dried and concentrated to a gum. The product was dissolved in dichloromethane and treated with 1.0M ethereal hydrogen chloride solution to give the titled product as a solid (0.1 g).

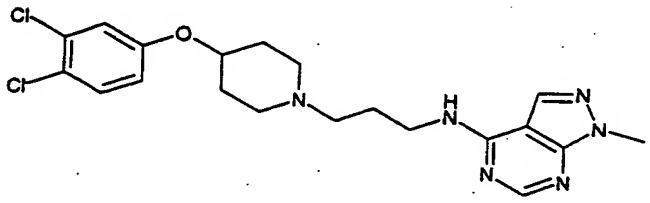
Melting point: 175-176 °C

MS: APCI(+ve): 389(M+H)

^{1}H NMR: δ(DMSO) 8.87 (t, 1H), 7.5 (m, 2H), 7.42 (m, 1H), 7.32 (m, 1H), 7.13 (m, 2H), 6.98 (m, 2H), 4.82 (m, 1/2H), 4.61 (m,1/2H), 3.81 (s, 3H), 3.69 (m, 3H), 3.68 (m, 3H), 3.47 (m, 1H), 3.13-3.22 (m, 4H), 2.27 (m, 1H), 2.14 (m, 1H), 2.03 (m, 1H), 1.90 (m, 1H)

Example 211

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine



(i) 2-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1H-isoindole-1,3(2H)-dione

A solution of the product from Example 1 step (ii) (2.0g), 2-(3-bromopropyl)-1H-isoindole-1,3(2H)-dione (1.61g) and triethylamine (2.5ml) in dichloromethane (40ml) was heated under reflux for 48h. The reaction mixture was partitioned between ethyl

acetate/water, the organic layer dried and evaporated under reduced pressure. Purification was by chromatography eluting with 4% methanol/dichloromethane. Yield 0.839g

MS: APCI(+ve) 433 (M+1)

(ii) 3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propylamine, dihydrochloride salt

The product from step (i) (0.83g) and hydrazine hydrate (0.1ml) in ethanol was heated under reflux for 6h. The precipitate was filtered off and partitioned between 2M hydrochloric acid and dichloromethane, the solid was filtered off and the aqueous layer basified with aqueous potassium hydroxide solution and extracted with dichloromethane. The organic layer was dried, evaporated under reduced pressure and the dihydrochloride salt formed using ethereal hydrogen chloride. Yield 0.28g

^{1}H NMR: δ(DMSO-d_6) 11.11(br s, 1H), 8.13(br s, 3H), 7.56 (d, 1H), 7.37(s, 1H), 7.10-7.06(br m, 1H), 4.84(br s, 0.5H), 4.65(br s,0.5H), 3.60-2.90(m, 8H), 2.24-2.01(m, 6H).

(iii) N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine

The product from step (ii) (0.08g), 4-chloro-1-methyl-1H-pyrazolo[3,4-d]pyrimidine (0.054g) and diisopropylethylamine (0.082g) in 1-methyl-2-pyrrolidinone (2ml) was heated at 50°C for 3h. The reaction mixture was diluted with ethyl acetate and washed with water. The organic layer was dried and the solvent removed under reduced pressure. Purification was by chromatography eluting with 9% methanol/dichloromethane. Yield 0.052g

MS: APCI(+ve) 435 (M+1)

^{1}H NMR: δ(DMSO-d_6) 8.25-8.22(m, 2H), 8.07(s, 1H), 7.49(d, 1H), 7.25(d, 1H), 6.97(dd, 1H), 4.46-4.42(m, 1H), 3.88(s, 3H), 3.49(q, 2H), 2.70-2.66(m, 2H), 2.40-2.36(m, 2H), 2.27-2.22(m, 2H), 1.92-1.88(m, 2H), 1.81-1.74(m, 2H), 1.62-1.59(m, 2H).

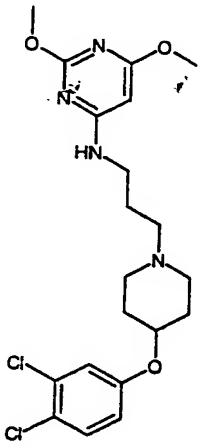
Melting point:120-124°C

Examples 212-255

The product from Example 211 step (ii) (1.5mg), the appropriate activated halo aromatic (1.25 equivalents), diisopropylethylamine (10 equivalents) in 1-methyl-2-pyrrolidinone (0.15ml) were heated at 100°C for 24h. The reaction mixture was evaporated to dryness and the residue dissolved in dimethylsulphoxide (0.4ml).

Example 212

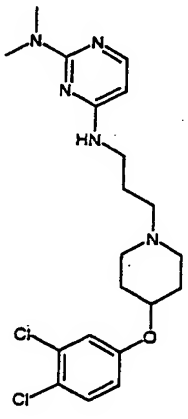
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2,6-dimethoxy-4-pyrimidinamine



MS: APCI(+ve) 441 (M+1)

Example 213

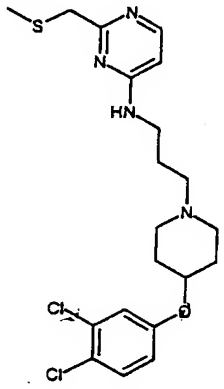
N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 424 (M+1)

Example 214

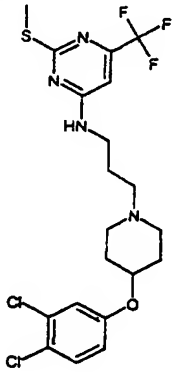
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine



MS: APCI(+ve) 441 (M+1)

Example 215

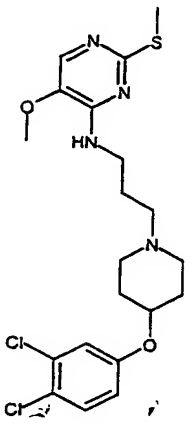
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-6-(trifluoromethyl)-4-pyrimidinamine



MS: APCI(+ve) 495 (M+1)

Example 216

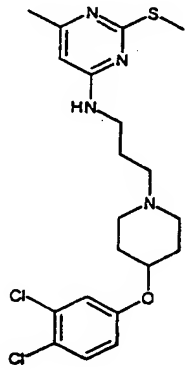
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 457 (M+1)

Example 217

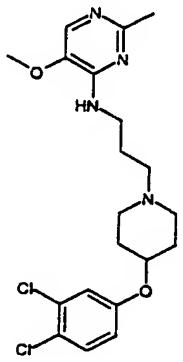
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 441 (M+1)

Example 218

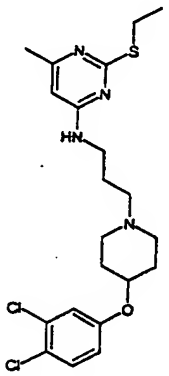
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-methyl-4-pyrimidinamine



MS: APCI(+ve) 425 (M+1)

Example 219

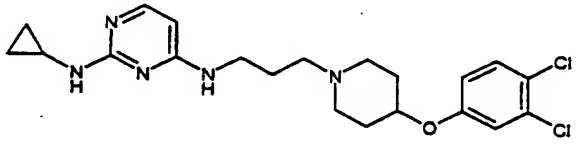
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(ethylsulfanyl)-6-methyl-4-pyrimidinamine



MS: APCI(+ve) 455 (M+1)

Example 220

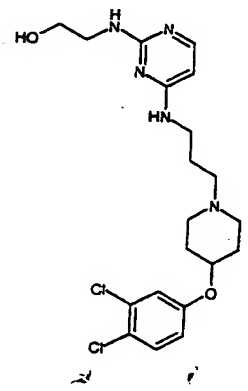
N~2~-Cyclopropyl-N~4~-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2,4-pyrimidinediamine



MS: APCI(+ve) 436 (M+1)

Example 221

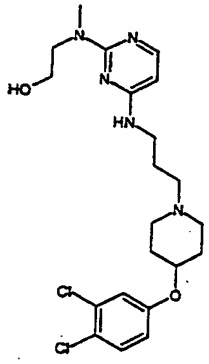
2-{[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl]amino}-1-ethanol



MS: APCI(+ve) 440 (M+1)

Example 222

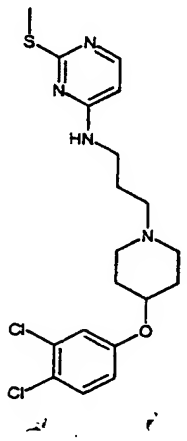
2-[[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol



MS: APCI(+ve) 454 (M+1)

Example 223

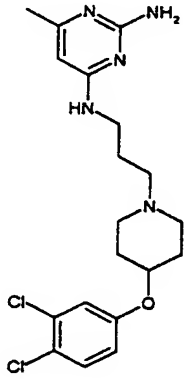
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 427 (M+1)

Example 224

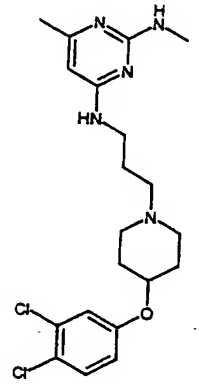
N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2,4-pyrimidinediamine



MS: APCI(+ve) 410 (M+1)

Example 225

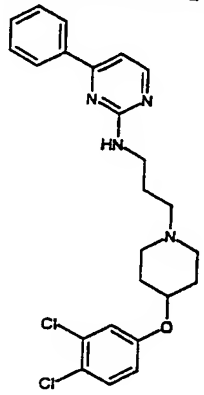
N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 424 (M+1)

Example 226

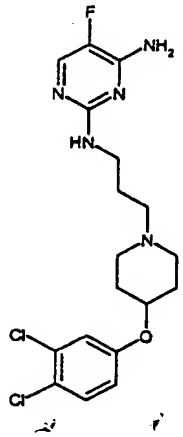
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-phenyl-2-pyrimidinamine



MS: APCI(+ve) 457 (M+1)

Example 227

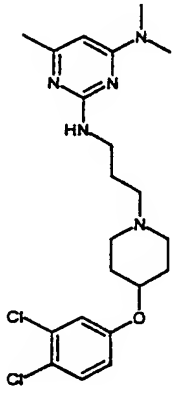
N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-fluoro-2,4-pyrimidinediamine



MS: APCI(+ve) 414 (M+1)

Example 228

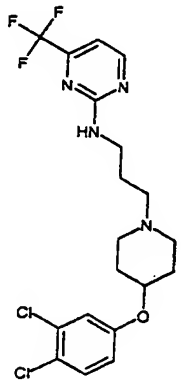
N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 438 (M+1)

Example 229

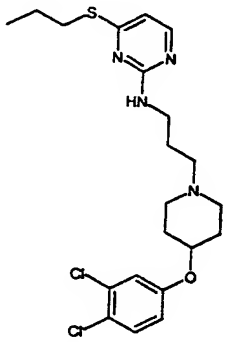
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(trifluoromethyl)-2-pyrimidinamine



MS: APCI(+ve) 449 (M+1)

Example 230

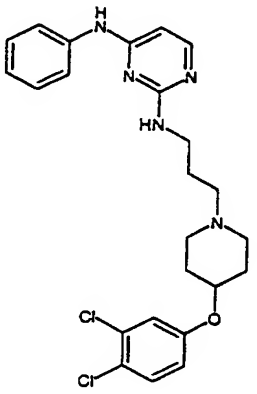
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(propylsulfanyl)-2-pyrimidinamine



MS: APCI(+ve) 455 (M+1)

Example 231

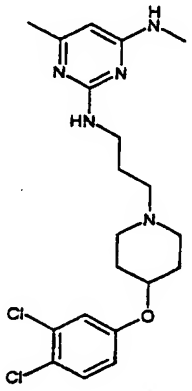
N~2~~{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~~phenyl-2,4-pyrimidinediamine



MS: APCI(+ve) 472 (M+1)

Example 232

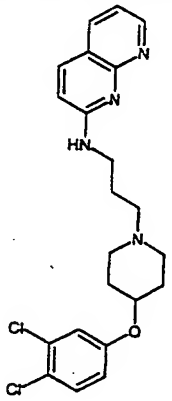
N~2~~{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,6-dimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 424 (M+1)

Example 233

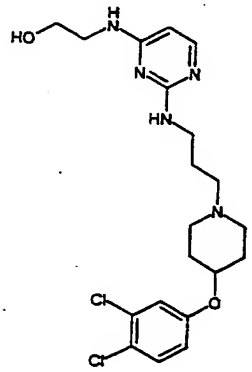
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}[1,8]naphthyridin-2-amine



MS: APCI(+ve) 431 (M+1)

Example 234

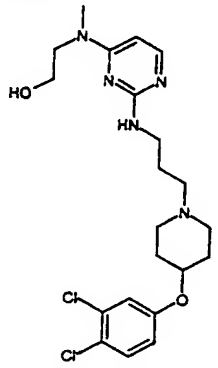
2-{[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl]amino}-1-ethanol



MS: APCI(+ve) 440 (M+1)

Example 235

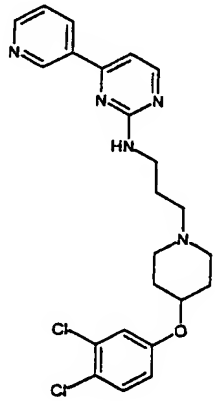
2-[[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl](methyl)amino]-1-ethanol



MS: APCI(+ve) 454 (M+1)

Example 236

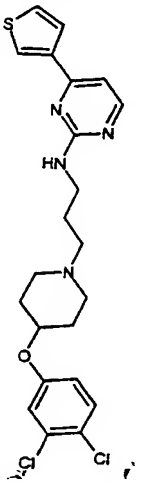
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-pyridinyl)-2-pyrimidinamine



MS: APCI(+ve) 458 (M+1)

Example 237

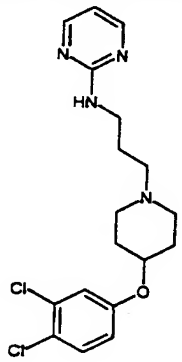
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-thienyl)-2-pyrimidinamine



MS: APCI(+ve) 463 (M+1)

Example 238

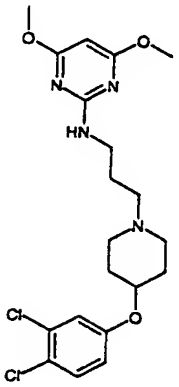
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrimidinamine



MS: APCI(+ve) 381 (M+1)

Example 239

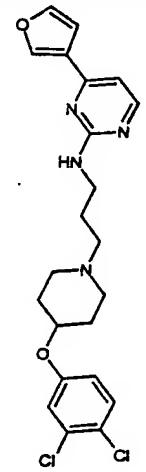
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4,6-dimethoxy-2-pyrimidinamine



MS: APCI(+ve) 441 (M+1)

Example 240

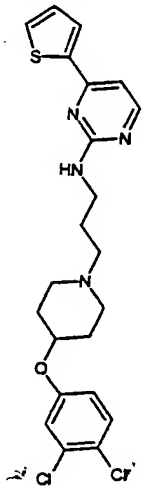
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-furyl)-2-pyrimidinamine



MS: APCI(+ve) 447 (M+1)

Example 241

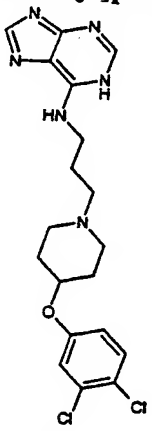
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(2-thienyl)-2-pyrimidinamine



MS: APCI(+ve) 463 (M+1)

Example 242

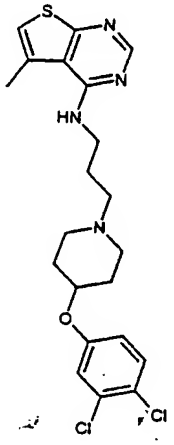
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1H-purin-6-amine



MS: APCI(+ve) 421 (M+1)

Example 243

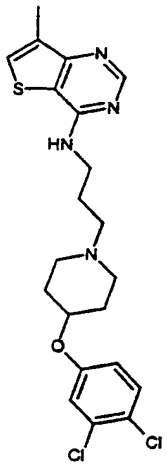
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methylthieno[2,3-d]pyrimidin-4-amine



MS: APCI(+ve) 451 (M+1)

Example 244

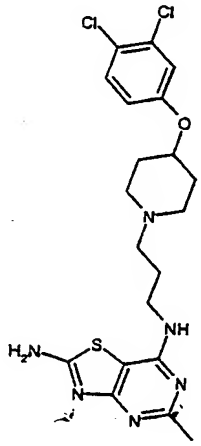
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-7-methylthieno[3,2-d]pyrimidin-4-amine



MS: APCI(+ve) 451 (M+1)

Example 245

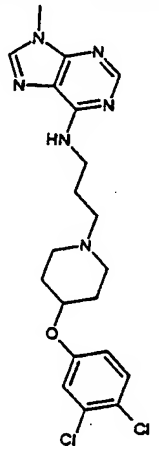
N~7~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine



MS: APCI(+ve) 467 (M+1)

Example 246

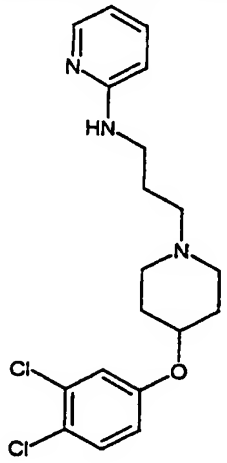
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-9-methyl-9H-purin-6-amine



MS: APCI(+ve) 435 (M+1)

Example 247

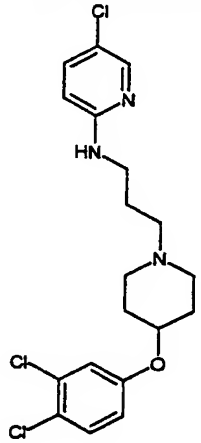
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine



MS: APCI(+ve) 379 (M+1)

Example 248

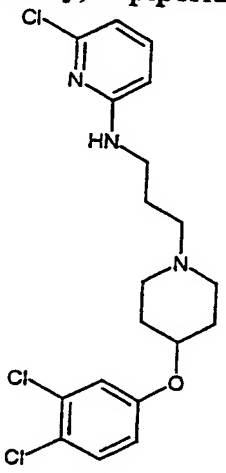
5-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine



MS: APCI(+ve) 414 (M+1)

Example 249

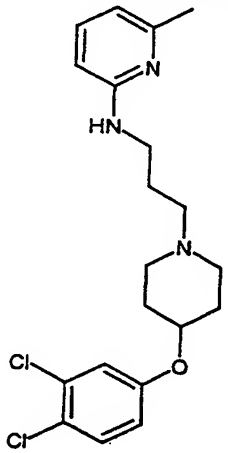
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine



MS: APCI(+ve) 414 (M+1)

Example 250

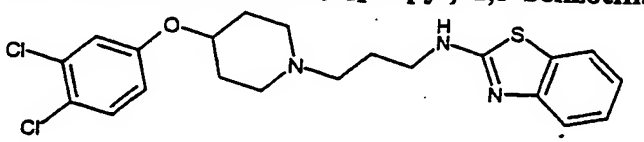
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-pyridinamine



MS: APCI(+ve) 494 (M+1)

Example 251

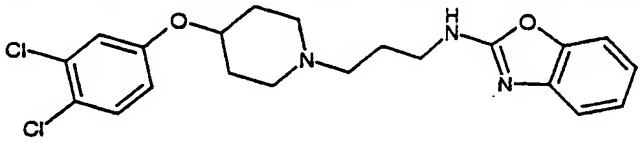
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzothiazol-2-amine



MS: APCI(+ve) 436 (M+1)

Example 252

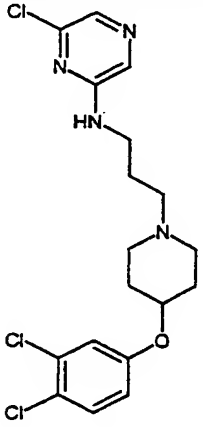
N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzoxazol-2-amine



MS: APCI(+ve) 420 (M+1)

Example 253

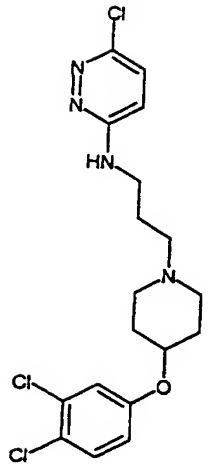
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrazinamine



MS: APCI(+ve) 415 (M+1)

Example 254

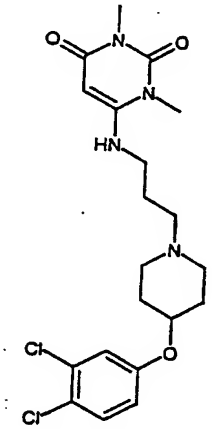
6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-3-pyridazinamine



MS: APCI(+ve) 417 (M+1)

Example 255

6-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione



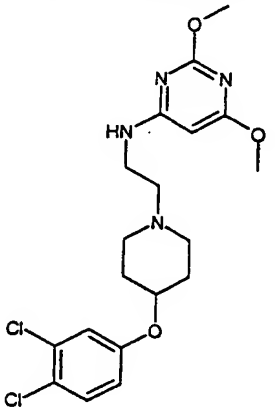
MS: APCI(+ve) 441 (M+1)

Examples 256-292

The product from Example 1 step (iv) (2.07mg), the appropriate activated halo aromatic (1.25 equivalents), diisopropylethylamine (10 equivalents) in 1-methyl-2-pyrrolidinone (0.15ml) were heated at 100°C for 24h.. The reaction mixture was evaporated to dryness and the residue dissolved in dimethylsulphoxide (0.4ml).

Example 256

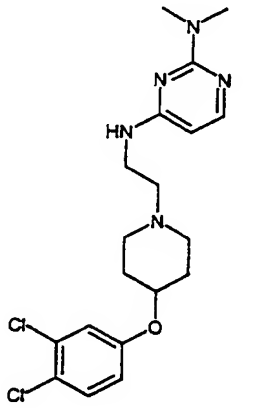
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,6-dimethoxy-4-pyrimidinamine



MS: APCI(+ve) 427 (M+1)

Example 257

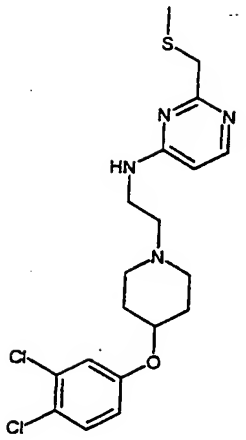
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine



MS: APCI(+ve) 410 (M+1)

Example 258

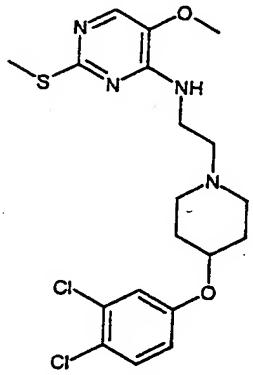
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine



MS: APCI(+ve) 427 (M+1)

Example 259

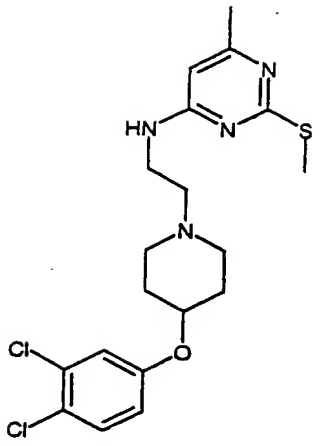
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 443 (M+1)

Example 260

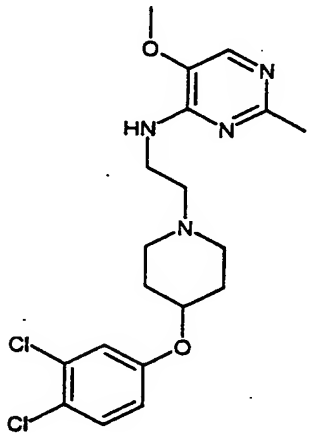
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine



MS: APCI(+ve) 427 (M+1)

Example 261

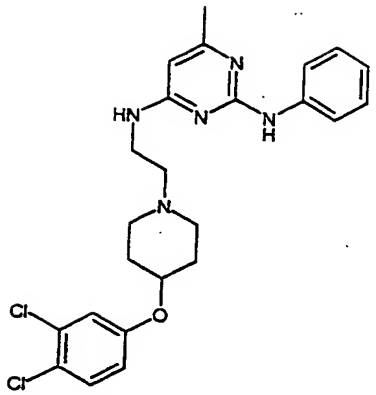
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-methyl-4-pyrimidinamine



MS: APCI(+ve) 411 (M+1)

Example 262

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-N~2~-phenyl-2,4-pyrimidinediamine



MS: APCI(+ve) 472 (M+1)

Pharmacological Analysis

Calcium flux $[Ca^{2+}]_i$ assay

a) Human eosinophils

Human eosinophils were isolated from EDTA anticoagulated peripheral blood as previously described (Hansel et al., *J. Immunol. Methods*, 1991, 145, 105-110). The cells were resuspended ($5x10^6$ ml^{-1}) and loaded with 5μM FLUO-3/AM + Pluronic F127 2.2μl/ml (Molecular Probes) in low potassium solution (LKS; NaCl 118mM, $MgSO_4$ 0.8mM, glucose 5.5mM, Na_2CO_3 8.5mM, KCl 5mM, HEPES 20mM, $CaCl_2$ 1.8mM, BSA 0.1%, pH 7.4) for one hour at room temperature. After loading, cells were centrifuged at 200g for 5min and resuspended in LKS at $2.5x10^6$ ml^{-1}. The cells were then transferred to 96 well FLIPr plates (Poly-D-Lysine plates from Becton Dickinson pre-incubated with 5μM fibronectin for two hours) at 100ml/well. The plate was centrifuged at 200g for 5min and the cells were washed twice with LKS (200μl; room temperature).

A compound of the Examples was pre-dissolved in DMSO and added to a final concentration of 0.1%(v/v) DMSO. Assays were initiated by the addition of an A_{50} concentration of eotaxin and the transient increase in fluo-3 fluorescence (l_{Ex} =490nm and l_{Em} = 520nm) monitored using a FLIPR (Fluorometric Imaging Plate Reader, Molecular Devices, Sunnyvale, U.S.A.).

b) **Human monocytes**

Human monocytes were isolated from EDTA anticoagulated peripheral blood as previously described (Cunoosamy & Holbrook, J. Leukocyte Biology, 1998, S2, 13). Cells were resuspended ($5x10^6$ ml^{-1}) in LKS and loaded with 5μM FLUO-3/AM + Pluronic F127 2.2μl/ml (Molecular Probes) for one hour at room temperature. After loading, cells were centrifuged at 200g for 5min and resuspended in LKS at $0.5x10^6$ ml^{-1}. The cells were then transferred to 96 well FLIPr plates (Costar). To each well 100μl of cells were added at a concentration of $0.5x10^6$ ml^{-1}. The plates were centrifuged (200g; 5 mins; room temperature) to allow the cells to adhere. After centrifugation the cells were washed twice with LKS (200μl; room temperature).

A compound of the Examples was pre-dissolved in DMSO and added to a final concentration of 0.1%(v/v) DMSO. Assays were initiated by the addition of an A_{50} concentration of MIP-1α and the transient increase in fluo-3 fluorescence (l_{Ex} =490nm and l_{Em} = 520nm) monitored using a FLIPR (Fluorometric Imaging Plate Reader, Molecular Devices, Sunnyvale, U.S.A.).

The compounds of the Examples were found to be antagonists of the eotaxin mediated $[Ca^{2+}]_i$ in human eosinophils and/or antagonists of the MIP-1α mediated $[Ca^{2+}]_i$ in human monocytes.

Human eosinophil chemotaxis

Human eosinophils were isolated from EDTA anticoagulated peripheral blood as previously described (Hansel et al., *J. Immunol. Methods*, 1991, 145, 105-110). The cells were resuspended at $10x10^6$ ml^{-1} in RPMI containing 200 IU/ml penicillin, 200 μg/ml streptomycin sulphate and supplemented with 10% HIFCS, at room temperature.

Eosinophils (700 μl) were pre-incubated for 15 mins at 37° C with 7 μl of either vehicle or compound (100x required final concentration in 10% DMSO). The chemotaxis plate

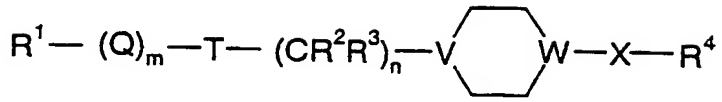
(ChemoTx, 3μm pore, Neuroprobe) was loaded by adding 28μl of a concentration of eotaxin (0.1 to 100nM) containing a concentration of a compound according to the Examples or solvent to the lower wells of the chemotaxis plate. The filter was then placed over the wells and 25 μl of eosinophil suspension were added to the top of the filter. The plate was incubated for 1 hr at 37° C in a humidified incubator with a 95% air/5% CO_2 atmosphere to allow chemotaxis.

The medium, containing cells that had not migrated, was carefully aspirated from above the filter and discarded. The filter was washed once with phosphate buffered saline (PBS) containing 5 mM EDTA to remove any adherent cells. Cells that had migrated through the filter were pelleted by centrifugation (300xg for 5 mins at room temperature) and the filter removed and the supernatant transferred to each well of a 96-well plate (Costar). The pelleted cells were lysed by the addition of 28 μl of PBS containing 0.5% Triton x100 followed by two cycles of freeze/thawing. The cell lysate was then added to the supernatant. The number of eosinophils migrating was quantified according to the method of Strath et al., *J. Immunol. Methods*, 1985, 83, 209 by measuring eosinophil peroxidase activity in the supernatant.

Certain compounds of the Examples were found to be antagonists of the eotaxin mediated human eosinophil chemotaxis.

CLAIMS

1. A compound of general formula



(I)

wherein:

R^1 represents a C_1-C_{12} alkyl group optionally substituted by one or more substituents independently selected from cyano, hydroxyl, C_1-C_6 alkoxy, C_1-C_6 alkylthio and C_1-C_6 alkoxycarbonyl groups, or

R^1 represents a 3- to 10-membered saturated or unsaturated ring system which may comprise up to two ring carbon atoms that form carbonyl groups and which may comprise up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur, the ring system being optionally substituted by one or more substituents independently selected from halogen atoms, and cyano, nitro, hydroxyl, C_1-C_6 alkyl, C_3-C_6 cycloalkyl, C_1-C_6 alkoxy, C_1-C_6 alkoxycarbonyl, C_1-C_6 haloalkyl, C_1-C_6 haloalkoxy, $-NR^5R^6$, C_3-C_6 cycloalkylamino, C_1-C_6 alkylthio, C_1-C_6 alkylthioC_1-C_6 alkyl, C_1-C_6 alkylcarbonylamino, $-C(O)NR^7R^8$, sulphonamido, (di)C_1-C_6 alkylsulphonamido, phenyl, phenylamino, nitrophenyl, pyridyl, pyridylthio, benzodioxanyl, thienyl, furanyl, and $C(O)R^9$-substituted C_1-C_6 alkyl or C_1-C_6 alkoxy groups;

m is 0 or 1;

Q represents a group OCH_2, C_1-C_4 alkylene or C_2-C_4 alkenylene;

T represents a group C(O)NH, or when m is 0, T may additionally represent a bond or a group NH, or when m is 1 and Q represents C_1-C_4 alkylene, T may additionally represent a group NH;

n is 1, 2, 3 or 4;

each R^2 independently represents a hydrogen atom or a C_1-C_4 alkyl group;

each R^3 independently represents a hydrogen atom or a C_1-C_4 alkyl group;

V represents a nitrogen atom;

W represents a nitrogen atom or a group CH;

X represents an oxygen atom or a group C(O), CH(OH), NH or N(C_1-C_6 alkyl), provided that when W represents a nitrogen atom, then X represents C(O);

R^4 represents a phenyl group optionally substituted by one or more substituents independently selected from halogen atoms, and amino, nitro, cyano, sulphonyl, sulphonamido, C_1-C_6 alkyl, C_1-C_6 haloalkyl, C_1-C_6 haloalkoxy and C_1-C_6 alkylsulphonyl groups;

R^5 and R^6 each independently represent a hydrogen atom or a C_1-C_6 alkyl or hydroxyC_1-C_6 alkyl group, or R^5 and R^6 together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring;

R^7 and R^8 each independently represent a hydrogen atom or a C_1-C_6 alkyl group; and

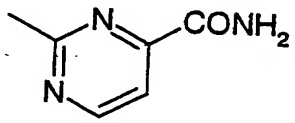
R^9 represents a hydroxyl or -NR^7R^8 group;

with the provisos that

(a) when m is 0, T is CONH, n is 2, 3 or 4 and each R^2 and R^3 represents hydrogen, W is CH, X is C(O) or CH(OH) and R^1 represents a substituted 3- to 10-membered unsaturated ring system, then the one or more substituents in the ring system do not include hydroxyl, halogen, C_1-C_6 alkoxy or C_1-C_6 haloalkoxy, and

(b) when W is N, X is C(O), R^4 represents 3-trifluoromethylphenyl, m is 0 and T is a bond, then R^1 and $(CR^2R^3)_n$ taken together do not represent a C_1-C_6 alkyl group, and

(c) when W is CH, X is O, n is 2 or 3 and each R^2 and R^3 represents hydrogen, m is 0 and T is NH, then R^1 does not represent a group



;

or a pharmaceutically acceptable salt or solvate thereof.

2. A compound according to claim 1, wherein R^1 represents a C_1-C_{10} alkyl group optionally substituted by one or two substituents independently selected from cyano, hydroxyl, C_1-C_4 alkoxy, C_1-C_4 alkylthio and C_1-C_4 alkoxycarbonyl groups, or

R^1 represents a 3- to 10-membered saturated or unsaturated ring system comprising up to two ring carbon atoms that form carbonyl groups and comprising up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur, the ring system being optionally substituted by one, two or three substituents independently selected from halogen atoms, and cyano, nitro, hydroxyl, C_1-C_4 alkyl, C_3-C_6 cycloalkyl, C_1-C_4 alkoxy, C_1-C_4 alkoxycarbonyl, C_1-C_3 haloalkyl, C_1-C_3 haloalkoxy, $-NR^5R^6$, C_3-C_6 cycloalkylamino, C_1-C_4 alkylthio, C_1-C_4 alkylthioC_1-C_4 alkyl, C_1-C_4 alkylcarbonylamino, $-C(O)NR^7R^8$, phenyl, phenylamino, nitrophenyl, pyridyl, pyridylthio, benzodioxanyl, thienyl, furanyl, and $C(O)R^9$-substituted C_1-C_4 alkyl or C_1-C_4 alkoxy groups.

3. A compound according to claim 1 or claim 2, wherein m is 1 and Q represents a group OCH_2, C_1-C_3 alkylene or C_2-C_3 alkenylene.

4. A compound according to any one of claims 1 to 3, wherein T represents a group C(O)NH.

5. A compound according to any one of the preceding claims, wherein n is 2 or 3.

6. A compound according to any one of the preceding claims, wherein V represents a nitrogen atom and W represents a group CH.

7. A compound according to any one of the preceding claims, wherein X represents an oxygen atom or a group C(O) or NH.

8. A compound according to any one of the preceding claims, wherein R^4 represents a phenyl group optionally substituted by one or two halogen atoms.

9. A compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, according to claim 1 being selected from:

4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-ethoxybenzamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-isopropoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-ethoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethoxy)benzamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide hydrochloride,

3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide hydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-hydroxybenzamide hydrochloride,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-[2-(methylamino)-2-oxoethoxy]benzamide hydrochloride,

2-[3-{2-[4-(4-Fluorobenzoyl)-1-piperidinyl]ethyl}-2,4-dioxo-3,4-dihydro-1(2H)-quinazolinyl]-N,N-dimethylacetamide hydrochloride,

N-{2-[4-(3,4-Dichlorobenzoyl)-1-piperazinyl]ethyl}-3-methoxybenzamide hydrochloride,

3,4-Dichloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}benzamide,

4-Chloro-N-{2-[4-(3,4-dichlorobenzoyl)-1-piperazinyl]ethyl}-2-[2-(dimethylamino)-2-oxoethoxy]benzamide hydrochloride,

N~7~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-9-methyl-9H-purin-6-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzothiazol-2-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzoxazol-2-amine,

6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrazinamine,

6-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-pyridazinamine,

6-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione,

N-{1-[4-(3,4-Dichlorophenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-4-methyl-benzamide, hydrochloride salt,

N-{1-[4-(3,4-Dichloro-phenoxy)-piperidin-1-ylmethyl]-2-methyl-propyl}-3-methoxy-benzamide, hydrochloride salt,

N-{2-[4-(3,4-Dichloroanilino)-1-piperidinyl]ethyl}-3-methoxybenzamide dihydrochloride,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N-(3-methoxybenzyl)amine dihydrochloride,

3-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methoxy-2,4(1H,3H)-quinazolinedione,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluorobenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-nitrobenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methylbenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-dinitrobenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-iodobenzamide,

4-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

4-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methylbenzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-nitrobenzamide,

3-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

3,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-fluorobenzamide,
2,4-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methylbenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-iodobenzamide,
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-nitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-5-(trifluoromethyl)benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(trifluoromethoxy)benzamide,
3,5-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)benzamide,
3-Cyano-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
2-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide,
3-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
2-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3,5-difluorobenzamide,
2,3-Dichloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-naphthamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-fluoro-6-(trifluoromethyl)benzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-difluorobenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-thiophenecarboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide,
Methyl 4-({2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}amino)-4-oxobutanoate,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclobutanecarboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-methoxyacetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclohexanecarboxamide,

(E)-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide,

2-Chloro-N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}nicotinamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenylacetamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}cyclopentanecarboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-phenoxyacetamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(trifluoromethyl)benzamide,

4-(tert-Butyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}benzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methylbenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-methylbenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-4-methyl-3-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-cyanobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-furamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-nitrobenzamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-naphthamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(methylsulfanyl)nicotinamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-(2,3-dihydro-1,4-benzodioxin-2-yl)-1,3-thiazole-4-carboxamide,

N~2~-Cyclopropyl-N~4~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine,

2-{[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl]amino}-1-ethanol,

2-[[4-({2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~-phenyl-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(methylsulfanyl)-4-pyrimidinamine,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2,4-pyrimidinediamine,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine,

2-Chloro-N~4~-cyclopropyl-N~6~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenyl-2-pyrimidinamine,

N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(trifluoromethyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(propylsulfanyl)-2-pyrimidinamine,

N~2~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~4~-phenyl-2,4-pyrimidinediamine,

N~4~-Cyclopropyl-N~2~-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}[1,8]naphthyridin-2-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-pyridinyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,6-dimethoxy-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(3-furyl)-2-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-purin-6-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methylthieno[2,3-d]pyrimidin-4-amine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methylthieno[3,2-d]pyrimidin-4-amine,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-thiophenecarboxamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-quinoxalinecarboxamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}bicyclo[2.2.1]hept-5-ene-2-carboxamide,

N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}cyclohexanecarboxamide,

(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-phenyl-2-propenamide,
N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-2-phenoxyacetamide,
(E)-N-{2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}-3-(4-nitrophenyl)-2-propenamide,
2-(1-Adamantyl)-N-{2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}acetamide,
(4-Chlorophenyl)(1-{2-[(2-fluoro-4,5-dimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3,4,5-trimethoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl){1-[2-(isobutylamino)ethyl]-4-piperidinyl}methanone,
4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-ethylhexanenitrile,
(4-Chlorophenyl)(1-{2-[(7-hydroxy-3,7-dimethyloctyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)[1-(2-{[(6-nitro-1,3-benzodioxol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,
[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,
(4-Chlorophenyl)[1-(2-{[3-nitro-4-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl)[1-(2-{[(E)-3-(4-nitrophenyl)-2-propenyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl){1-[2-({[5-(3-nitrophenyl)-2-furyl]methyl}amino)ethyl]-4-piperidinyl}methanone,
(4-Chlorophenyl)[1-(2-{[5-nitro-2-(2-pyridinylsulfanyl)benzyl]amino}ethyl)-4-piperidinyl]methanone,
6-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-2-(methylsulfanyl)nicotinonitrile,
{1-[2-({[5-Chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazol-4-yl]methyl}amino)ethyl]-4-piperidinyl}(4-chlorophenyl)methanone,
(4-Chlorophenyl)[1-(2-{[3-(methylsulfanyl)butyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(4-phenyl-4-piperidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(1-phenyl-1H-pyrazol-5-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,

Ethyl 3-[({2-[4-(4-chlorobenzoyl)-1-piperidinyl]ethyl}-amino)methyl]cyclohexanecarboxylate,

N-{4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]phenyl}acetamide,

(4-Chlorophenyl)(1-{2-[(2,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(2,6-dichlorobenzyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)(1-{2-[(2-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone,

(4-Chlorophenyl)[1-(2-{[(3-methyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-methoxybenzyl)amino]ethyl}-4-piperidinyl)methanone,

3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4H-chromen-4-one,

[1-(2-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,

(4-Chlorophenyl)[1-(2-{[(2,6-dichloro-4-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(2-phenyl-1H-imidazol-4-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(5-ethyl-2-thienyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(2-chloro-3-quinolinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)[1-(2-{[(6-methyl-2-pyridinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,

(4-Chlorophenyl)(1-{2-[(3-quinolinylmethyl)amino]ethyl}-4-piperidinyl)methanone,

4-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-1,5-dimethyl-2-phenyl-1,2-dihydro-3H-pyrazol-3-one,

(4-Chlorophenyl)(1-{2-[(4-pyridinylmethyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3-hydroxy-4-nitrobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(4-Chlorophenyl)(1-{2-[(3,5-difluorobenzyl)amino]ethyl}-4-piperidinyl)methanone,
(1-{2-[(2-Chloro-6-fluorobenzyl)amino]ethyl}-4-piperidinyl)(4-chlorophenyl)methanone,
[1-(2-{[(4-Bromo-1H-pyrazol-3-yl)methyl]amino}ethyl)-4-piperidinyl](4-chlorophenyl)methanone,
3-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-6,7-dimethyl-4H-chromen-4-one,
2-{2-[({2-[4-(4-Chlorobenzoyl)-1-piperidinyl]ethyl}amino)methyl]-4-nitrophenoxy}acetic acid,
(4-Chlorophenyl)[1-(2-{[(1-methyl-1H-benzimidazol-2-yl)methyl]amino}ethyl)-4-piperidinyl]methanone,
(4-Chlorophenyl)[1-(2-{[(2,4-dimethoxy-5-pyrimidinyl)methyl]amino}ethyl)-4-piperidinyl]methanone,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-(methylamino)benzamide,
4-Chloro-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxy-4-methylbenzamide,
3-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-4-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-benzodioxole-5-carboxamide,
4-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-fluoro-4-methoxybenzamide,
5-Bromo-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-methyl-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4,5-dimethyl-2-furamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-ethoxy-1-benzofuran-2-carboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-1-benzofuran-2-carboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-7-methoxy-1-benzofuran-2-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-fluorophenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methoxyphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methylphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(2-methylphenyl)acetamide,
2-(3-Bromophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,
2-(2-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,
2-(4-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[2-(trifluoromethyl)phenyl]acetamide,
2-(3-Chlorophenyl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dimethoxyphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methoxyphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-dichlorophenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-fluoro-4-methoxyphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-ethoxyphenyl)acetamide,
2-(1,3-Benzodioxol-5-yl)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[4-(dimethylamino)phenyl]acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-methylphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3,4-difluorophenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(3-methoxyphenyl)acetamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-4-phenylbutanamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-phenylpropanamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-3-(3-methoxyphenyl)propanamide,
2-Amino-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide,
2-(Acetylamino)-N-{2-[4-(3,4-dichlorophenoxy)-1-piperidinyl]ethyl}-1,3-thiazole-4-carboxamide,
N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-(4-pyridinyl)-1,3-thiazole-4-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,4-dimethyl-1,3-thiazole-5-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,5-dimethyl-1,3-oxazole-4-carboxamide,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-1H-imidazole-4-carboxamide,

N-{2-[4-(3,4-Chlorophenoxy)-1-piperidinyl]ethyl}-3-methoxybenzamide, hydrochloride salt,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2,6-dimethoxy-4-pyrimidinamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-6-(trifluoromethyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methoxy-2-methyl-4-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(ethylsulfanyl)-6-methyl-4-pyrimidinamine,

N~2~-Cyclopropyl-N~4~-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2,4-pyrimidinediamine,

2-{[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl]amino}-1-ethanol,

2-[[4-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-2-pyrimidinyl](methyl)amino]-1-ethanol,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-(methylsulfanyl)-4-pyrimidinamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2,4-pyrimidinediamine,

N~4~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~2~,6-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-phenyl-2-pyrimidinamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-fluoro-2,4-pyrimidinediamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,N~4~,6-trimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(trifluoromethyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(propylsulfanyl)-2-pyrimidinamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~-phenyl-2,4-pyrimidinediamine,

N~2~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-N~4~,6-dimethyl-2,4-pyrimidinediamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}[1,8]naphthyridin-2-amine,

2-{[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl]amino}-1-ethanol,

2-[[2-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-4-pyrimidinyl](methyl)amino]-1-ethanol,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-pyridinyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-thienyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4,6-dimethoxy-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(3-furyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-4-(2-thienyl)-2-pyrimidinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1H-purin-6-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methylthieno[2,3-d]pyrimidin-4-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-7-methylthieno[3,2-d]pyrimidin-4-amine,

N~7~-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-5-methyl[1,3]thiazolo[4,5-d]pyrimidine-2,7-diamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-9-methyl-9H-purin-6-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,

5-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,

6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyridinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-6-methyl-2-pyridinamine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzothiazol-2-amine,

N-{3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}-1,3-benzoxazol-2-amine,

6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-2-pyrazinamine,

6-Chloro-N-{3-[4-(3,4-dichlorophenoxy)-1-piperidinyl]propyl}-3-pyridazinamine,

6-({3-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]propyl}amino)-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2,6-dimethoxy-4-pyrimidinamine,

N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-N~2~,N~2~-dimethyl-2,4-pyrimidinediamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-2-[(methylsulfanyl)methyl]-4-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-(methylsulfanyl)-4-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-2-(methylsulfanyl)-4-pyrimidinamine,

N-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-5-methoxy-2-methyl-4-pyrimidinamine, and

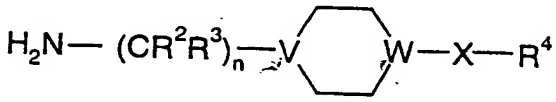
N~4~-{2-[4-(3,4-Dichlorophenoxy)-1-piperidinyl]ethyl}-6-methyl-N~2~-phenyl-2,4-pyrimidinediamine.

10. A process for the preparation of a compound of formula (I) as defined in claim 1 which comprises

(i) when T represents a group C(O)NH, reacting a compound of general formula

$$R^1\text{-}(Q)_m\text{-}COL^1 \quad (II)$$

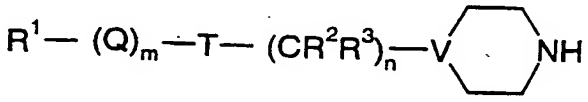
wherein L^1 represents a leaving group (e.g. a hydroxyl or halide, such as chloride, group) and R^1, m and Q are as defined in formula (I), with a compound of general formula



(III)

or an acid addition salt thereof (e.g. trifluoroacetate) wherein n, R^2, R^3, V, W, X and R^4 are as defined in formula (I); or

(ii) when T represents a group C(O)NH and W represents a nitrogen atom, reacting a compound of general formula



(IV)

wherein R^1, m, Q, T, n, R^2, R^3 and V are as defined in formula (I), with a compound of general formula

$$L^2\text{-}X\text{-}R^4 \quad (V)$$

wherein L^2 represents a leaving group (e.g. a halogen atom) and X and R^4 are as defined in formula (I); or

(iii) when T represents a group NH and m is 0, reacting a compound of general formula

$$R^1\text{-}L^3 \quad (VI)$$

wherein L^3 represents a leaving group (e.g. a halogen atom) and R^1 is as defined in formula (I), with a compound of formula (III) as defined in (i) above; or

(iv) when T represents a group NH, m is 1 and Q represents C_1-C_4 alkylene, reacting a compound of general formula

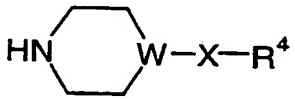
$$R^1\text{-}(CH_2)_p\text{-}CHO \quad \text{(VII)}$$

wherein p is 0, 1, 2 or 3 and R^1 is as defined in formula (I), with a compound of formula (III) as defined in (i) above; or

(v) when T represents a bond and m is 0, reacting a compound of general formula

$$R^1\text{-}(CR^2R^3)_n\text{-}L^4 \quad \text{(VIII)}$$

wherein L^4 represents a leaving group such as a halogen atom (e.g. chlorine) and n, R^1, R^2 and R^3 are as defined in formula (I), with a compound of general formula



(IX)

wherein W, X and R^4 are as defined in formula (I);

and optionally after (i), (ii), (iii), (iv) or (v) converting the compound of formula (I) to a further compound of formula (I) and/or forming a pharmaceutically acceptable salt or solvate of the compound of formula (I).

11. A pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

12. A process for the preparation of a pharmaceutical composition as claimed in claim 11 which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9 with a pharmaceutically acceptable adjuvant, diluent or carrier.

13. A compound of formula (I), or a pharmaceutically-acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9 for use in therapy.

14. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9 in the manufacture of a medicament for use in therapy.

15. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9 in the manufacture of a medicament for the treatment of human diseases or conditions in which modulation of chemokine receptor activity is beneficial.

16. A method of treating an inflammatory disease in a patient suffering from, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 9.

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference F 2150-1 WO	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
International application No. PCT/SE 00/00563	International filing date *(day/month/year)* 22 March 2000	(Earliest) Priority Date *(day/month/year)* 26 March 1999
Applicant AstraZeneca UK Limited et al		

This international search report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This international search report consists of a total of 7 sheets.

[X] It is also accompanied by a copy of each prior art document cited in this report.

1. [X] **Certain claims were found unsearchable** (See Box I).

2. [ ] **Unity of invention is lacking** (See Box II).

3. [ ] The international application contains disclosure of a **nucleotide and/or amino acid sequence listing** and the international search was carried out on the basis of the sequence listing
   - [ ] filed with the international application.
   - [ ] furnished by the applicant separately from the international application,
     - [ ] but not accompanied by a statement to the effect that it did not include matter going beyond the disclosure in the international application as filed.
   - [ ] transcribed by this Authority.

4. With regard to the title,
   - [X] the text is approved as submitted by the applicant.
   - [ ] the text has been established by this Authority to read as follows:

5. With regard to the **abstract**,
   - [ ] the text is approved as submitted by the applicant.
   - [X] the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is:
   Figure No. ---
   - [ ] as suggested by the applicant.
   - [ ] because the applicant failed to suggest a figure.
   - [ ] because this figure better characterizes the invention.
   - [ ] None of the figures.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE00/00563

Box I **Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 16
because they relate to subject matter not required to be searched by this Authority, namely:

See PCT Rule 39.1.(iv).: Methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods.

2. ☒ Claims Nos.: 1-16
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

See extra sheet

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).:

Box II **Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE00/00563

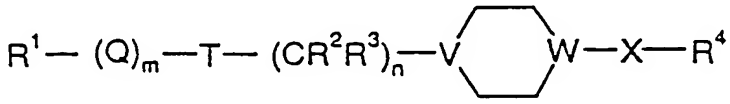
Present claims 1-16 relate to an extremely large number of possible compounds. In fact, the claim contains so many options, variables, possible permutations and provisions that a lack of clarity and conciseness within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible.

Consequently, the search has been carried out for those parts of the application which appear to be clear and concise, namely the examples and closely related homologous compounds etc., those mentioned in the description.

"The initial phase of the search revealed a very large number of documents relevant to the issue of novelty. So many documents were retrieved that it is impossible to determine which parts of the claim(s) may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons, a meaningful search over the whole breadth of the claims(s) is impossible. Consequently, the search has been restricted.

Box III TEXT OF THE ABSTRACT (Continuation of item 5 of the first sheet)

The invention provides compounds of general formula



(I)

wherein:

R^1 represents optionally substituted C_1-C_{12} alkyl or optionally substituted 3- to 10-membered saturated or unsaturated ring system comprising up to two ring carbon atoms that form carbonyl groups and comprising up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulfur;

m is 0-1;

Q represents OCH_2, C_1-C_4 alkylene or C_2-C_4 alkenylene;

T represents C(O)NH, or when m is 0, T may additionally represent a bond or NH, or when m is 1 and Q represents C_1-C_4 alkylene, T may additionally represent NH;

n is 1-4;

each R^2 and R^3 independently represents H or C_1-C_4 alkyl;

V represents N, and W represents N or CH;

X represents O, C(O), CH(OH), SO_2, NH or N(C_1-C_6 alkyl), provided that when W represents N, then X represents either C(O) or SO_2 and when W represents CH, then X is other than SO_2;

R^4 represents optionally substituted phenyl;

R^5 and R^6 each independently represent H, C_1-C_6 alkyl or hydroxyC_1-C_6 alkyl, or R^5 and R^6 together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring;

R^7 and R^8 each independently represent H or C_1-C_6 alkyl; and

R^9 represents OH or $-NR^7R^8$;

processes for their preparation, pharmaceutical compositions containing them and their use in therapy.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 00/00563

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 401/12, C07D 401/06, C07D 211/14, C07D 211/22, A61K 31/445, A61P 11/00, A61P 19/00, A61P 31/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K, A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5210086 A (PASCAL, GEORGE ET AL), 11 May 1993 (11.05.93)	1-16
	--	
X	US 5143923 A (NICHOLAS J. HRIB ET AL), 1 Sept 1992 (01.09.92)	1-16
	--	
X	US 4853393 A (ZIMMERMANN, MARKUS), 1 August 1989 (01.08.89)	1-16
	--	
X	US 4335127 A (VANDERBERK JAN ET AL), 15 June 1982 (15.06.82)	1-16
	--	

[X] Further documents are listed in the continuation of Box C. [X] See patent family annex.

* Special categories of cited documents:
"A" document defining the general state of the art which is not considered to be of particular relevance
"E" erlier document but published on or after the international filing date
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"O" document referring to an oral disclosure, use, exhibition or other means
"P" document published prior to the international filing date but later than the priority date claimed
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
22 June 2000	04 -08- 2000
Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. +46 8 666 02 86	Authorized officer Eva Johansson/gh Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 00/00563

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9742956 A1 (SYNAPTIC PHARMACEUTICAL CORPORATION), 20 November 1997 (20.11.97)	1-16
X	EP 0288563 A1 (EISAI CO., LTD), 2 November 1988 (02.11.88)	1-16
X	EP 0184258 A2 (JANSSEN PHARMACEUTICA N.V.), 11 June 1986 (11.06.86)	1-16
X	EP 0124476 A1 (CIBA-GEIGY AG), 7 November 1984 (07.11.84)	1-16
X	FR 2724382 A1 (SYNTHELABO SOCIETE ANONYME), 15 March 1996 (15.03.96)	1-16
X	STN International, File CAPLUS, CAPLUS accession no. 1997:259827, Document no. 127:17595, Kyorin Pharmaceutical Co., Ltd.: "Preparation of benzamide derivatives as gastrointestinal movement modulators" & JP,A2,09077742,19970325	1-16
X	STN International, File CAPLUS, CAPLUS accession no. 1997:204127, Document no. 126:212045, Yamonuchi Pharma Co Ltd: "Preparation of fused benzene ring derivatives as antidepressant drugs"; & JP,A2,09040646,19970210	1-16
X	STN International, File CAPLUS, CAPLUS accession no. 1992:151788, Document no. 116:151788, Daiichi Seiyaku Co.,Ltd.: "Preparation of pyrimidine derivatives as serotonin 2 receptor antagonists with high selectivity"; & JP,A2,03264579,19911125	1-16

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 00/00563

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	J.Med.Chem., Volume 37, 1994, NicholasJ. Hrib et al, "Benzisoxazole- and Benzisothiazole-3-carboxamides as Potential Atypical Antipsychotic Agents" page 2308 - page 2314 --	1-16
X	J.Med.Chem., Volume 35, 1992, Jeff L. Herndon et al, "Ketanserin Analogues: Structure-Affinity Relationships for 5-HT2 and 5-HT1c serotonin receptor binding" page 4903 - page 4910 --	1-16
X	Journal of Labelled Compounds and Radiopharmaceuticals, Volume 25, No 7, November 1987, Cor G:M: Jansosn et al, "Synthesis of 3H and 14C ketaserin" page 783 - page 792 -- --------	1-16

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.
PCT/SE 00/00563

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5210086 A	11/05/93	AU 637261 B	20/05/93
		AU 8487191 A	09/04/92
		CA 2052609 A	03/04/92
		CS 9102999 A	15/04/92
		EP 0480794 A	15/04/92
		FI 914611 A	03/04/92
		FR 2667317 A,B	03/04/92
		IL 99612 D	00/00/00
		JP 4273870 A	30/09/92
		MX 9101341 A	05/06/92
		NO 913843 A	03/04/92
		NZ 240055 A	25/09/92
		PT 99138 A	31/12/93
US 5143923 A	01/09/92	AT 142628 T	15/09/96
		AU 644054 B	02/12/93
		AU 1518792 A	05/11/92
		CA 2067404 A	30/10/92
		CS 9201298 A	18/11/92
		CZ 282764 B	15/10/97
		DE 69213544 D,T	27/02/97
		DK 511610 T	16/06/97
		EP 0511610 A,B	04/11/92
		SE 0511610 T3	
		ES 2094843 T	01/02/97
		FI 921875 A	30/10/92
		GR 3021211 T	31/12/96
		HU 214032 B	29/12/97
		IE 77036 B	19/11/97
		IL 101700 A	27/11/95
		JP 2656189 B	24/09/97
		JP 5132472 A	28/05/93
		MX 9201963 A	01/11/92
		NO 180488 B,C	20/01/97
		NO 921654 D	00/00/00
		NZ 242479 A	24/02/95
		PL 168090 B	31/01/96
		PL 168870 B	30/04/96
		RU 2039057 C	09/07/95
		US 5225412 A	06/07/93

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.
PCT/SE 00/00563

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4853393 A	01/08/89	AU 4744985 A	20/03/86
		DD 237165 A	02/07/86
		DK 416885 A	15/03/86
		DK 416985 A	15/03/86
		EP 0174653 A	19/03/86
		EP 0174654 A	19/03/86
		ES 546945 A	16/11/86
		ES 546946 A	16/02/87
		ES 554487 A	01/07/87
		ES 554488 A	01/12/87
		FI 853477 A	15/03/86
		FI 853478 A	15/03/86
		GR 852215 A	14/01/86
		GR 852216 A	14/01/86
		IL 76366 A	31/07/89
		JP 61072751 A	14/04/86
		JP 61072752 A	14/04/86
		NO 853609 A	17/03/86
		NZ 213461 A	27/07/89
		PT 81116 A,B	01/10/85
		PT 81117 A,B	01/10/85

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.
PCT/SE 00/00563

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4335127 A	15/06/82	AT 5258 E,T	15/11/83
		AU 536175 B	19/04/84
		AU 5438180 A	17/07/80
		CA 1132557 A	28/09/82
		CS 223977 B	25/11/83
		CY 1252 A	31/08/84
		DK 7280 A	09/07/80
		DK 170669 B	27/11/95
		EP 0013612 A,B	23/07/80
		SE 0013612 T3	
		ES 487537 A	16/12/80
		FI 66609 B,C	31/07/84
		FI 800047 A	09/07/80
		GR 67304 A	29/06/81
		HK 76284 A	19/10/84
		HR 930474 B	31/08/97
		IE 49351 B	18/09/85
		IL 59084 A	29/02/84
		JP 1497429 C	29/05/89
		JP 55105679 A	13/08/80
		JP 63046753 B	19/09/88
		KR 8500683 B	14/05/85
		KR 8500684 B	14/05/85
		LU 88219 A	03/02/94
		MY 55587 A	31/12/87
		NO 800034 A	09/07/80
		NZ 192551 A	06/07/84
		PH 17114 A	01/06/84
		PT 70662 A	01/02/80
		RO 79148 A	17/08/82
		SG 48684 G	29/03/85
		SI 8010046 A	31/12/94
		US 4522945 A	11/06/85
		YU 4680 A	30/06/83
		YU 42957 B	28/02/89
		ZA 8000082 A	26/08/81
WO 9742956 A1	20/11/97	AU 3008297 A	05/12/97

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.
PCT/SE 00/00563

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0288563 A1	02/11/88	SE 0288563 T3	
		AT 105550 T	15/05/94
		AU 599339 B	19/07/90
		AU 6405486 A	21/04/88
		DE 3689836 D,T	22/09/94
		DK 273788 A	19/05/88
		FI 90533 B,C	15/11/93
		FI 882369 A	19/05/88
		NO 175055 B,C	16/05/94
		WO 8802365 A	07/04/88
EP 0184258 A2	11/06/86	SE 0184258 T3	
		AT 75741 T	15/05/92
		CA 1246074 A	06/12/88
		DE 3585990 A	11/06/92
		JP 2009087 C	11/01/96
		JP 7045492 B	17/05/95
		JP 61148175 A	05/07/86
		US 4665075 A	12/05/87
EP 0124476 A1	07/11/84	SE 0124476 T3	
		AT 35257 T	15/07/88
		AU 583470 B	04/05/89
		AU 2542484 A	13/09/84
		CA 1273929 A	11/09/90
		CY 1576 A	20/12/91
		DD 220308 A	27/03/85
		DE 3472251 A	28/07/88
		DK 114684 A	10/09/84
		DK 159420 B,C	15/10/90
		ES 544713 A	16/01/87
		ES 553050 A	16/10/87
		FI 80019 B,C	29/12/89
		FI 840867 A	10/09/84
		HK 18491 A	22/03/91
		IE 57057 B	08/04/92
		IL 71185 A	30/06/88
		JP 1844916 C	25/05/94
		JP 59186959 A	23/10/84
		NO 840889 A	10/09/84
		NZ 207428 A	06/03/87
		PH 21416 A	15/10/87
		PT 78212 A,B	01/04/84
		SG 91490 G	18/01/91
		US 4559349 A	17/12/85
		ZA 8401728 A	28/11/84
		GR 82643 A	07/02/85
FR 2724382 A1	15/03/96	NONE	